

DECLARATION

I, Jane Roberta Mann, B.A., a Translator, of Frank B. Dehn & Co., 59 St Aldates, Oxford OX1 1ST, England, do declare that I have a competent knowledge of the English and German languages and that the document that is annexed hereto is a true and accurate translation of the German text of the U.S. Provisional Application Serial No. 60/408,021 filed September 04, 2002.

I further declare that all statements made of my own knowledge are true and that all statements made on information and belief are believed to be true.

I acknowledge that wilful false statements and the like are punishable by fine or imprisonment, or both [18 U.S.C. 1001] and may jeopardize the validity of the application or any patent issuing therefrom.

A handwritten signature in cursive script, reading "Jane Mann", is written over a horizontal line.

Signed this 15th day of July, 2003

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Boehringer Ingelheim Pharma KG

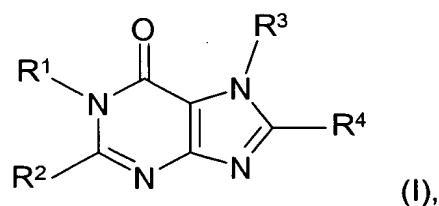
D-55216 Ingelheim/Rhein

Case 1/1385

Priority text

New purine derivatives, the preparation thereof and their use as
pharmaceutical compositions

The present invention relates to substituted purines of general formula



the tautomers, the stereoisomers, the mixtures, the prodrugs thereof and the salts thereof, particularly the physiologically acceptable salts thereof with inorganic or organic acids or bases which have valuable pharmacological properties, particularly an inhibiting effect on the activity of the enzyme dipeptidylpeptidase-IV (DPP-IV), the preparation thereof, the use thereof for the prevention or treatment of diseases or conditions associated with an increased DPP-IV activity or capable of being prevented or alleviated by reducing the DPP-IV activity, particularly type I or type II diabetes mellitus, the pharmaceutical compositions containing a compound of general formula (I) or a physiologically acceptable salt thereof as well as processes for the preparation thereof.

In the above formula I

R¹ denotes a hydrogen atom,

a C₁₋₈-alkyl group,

a C₃₋₈-alkenyl group,

a C₃₋₄-alkenyl group which is substituted by a C₁₋₂-alkyloxy-carbonyl, aminocarbonyl, C₁₋₃-alkylamino-carbonyl, di-(C₁₋₃-alkyl)-amino-carbonyl, pyrrolidin-1-ylcarbonyl, piperidin-1-ylcarbonyl or morpholin-4-ylcarbonyl group,

a C₃₋₈-alkynyl group,

a C₁₋₆-alkyl group substituted by a group R_a, where

R_a denotes a C₃₋₇-cycloalkyl, heteroaryl, cyano, carboxy, C₁₋₃-alkyloxy-carbonyl, aminocarbonyl, C₁₋₃-alkylamino-carbonyl, di-(C₁₋₃-alkyl)-amino-carbonyl, pyrrolidin-1-ylcarbonyl, piperidin-1-ylcarbonyl, morpholin-4-ylcarbonyl, piperazin-1-ylcarbonyl, 4-methylpiperazin-1-ylcarbonyl or 4-ethylpiperazin-1-ylcarbonyl group,

a C₁₋₆-alkyl group substituted by a phenyl group, where the phenyl ring is substituted by the groups R¹⁰ to R¹⁴ and

R¹⁰ denotes a hydrogen atom,

a fluorine, chlorine, bromine or iodine atom,

a C₁₋₄-alkyl, hydroxy or C₁₋₄-alkyloxy group,

a nitro, amino, C₁₋₃-alkylamino, di-(C₁₋₃-alkyl)-amino, cyan-C₁₋₃-alkylamino, *N*-(cyan-C₁₋₃-alkyl)-*N*-(C₁₋₃-alkyl)-amino, C₁₋₃-alkyloxy-carbonyl-C₁₋₃-alkylamino, pyrrolidin-1-yl, piperidin-1-yl, morpholin-4-yl, piperazin-1-yl or 4-(C₁₋₃-alkyl)-piperazin-1-yl group,

a formylamino, C₁₋₃-alkyl-carbonylamino, C₃₋₆-cycloalkyl-carbonylamino, C₃₋₆-cycloalkyl-C₁₋₃-alkyl-carbonylamino, arylcarbonylamino, aryl-C₁₋₃-alkyl-carbonylamino, C₁₋₃-alkyloxy-carbonylamino, aminocarbonylamino, C₁₋₃-alkyl-aminocarbonylamino, di-(C₁₋₃-alkyl)-aminocarbonylamino, pyrrolidin-1-yl-carbonylamino, piperidin-1-yl-carbonylamino, morpholin-4-

yl-carbonylamino, piperazin-1-yl-carbonylamino or 4-(C₁₋₃-alkyl)-piperazin-1-yl-carbonylamino, C₁₋₃-alkyl-sulphonylamino, bis-(C₁₋₃-alkylsulphonyl)-amino, aminosulphonylamino, C₁₋₃-alkylamino-sulphonylamino, di-(C₁₋₃-alkyl)-amino-sulphonylamino, pyrrolidin-1-yl-sulphonylamino, piperidin-1-yl-sulphonylamino, morpholin-4-yl-sulphonylamino, piperazin-1-yl-sulphonylamino or 4-(C₁₋₃-alkyl)-piperazin-1-yl-sulphonylamino, (C₁₋₃-alkylamino)-thiocarbonylamino, (C₁₋₃-alkyloxy-carbonylamino)-carbonylamino, arylsulphonylamino or aryl-C₁₋₃-alkyl-sulphonylamino group,

an N-(C₁₋₃-alkyl)-formylamino, *N*-(C₁₋₃-alkyl)-*N*-(C₁₋₃-alkyl-carbonyl)-amino, *N*-(C₁₋₃-alkyl)-*N*-(C₃₋₆-cycloalkyl-carbonyl)-amino, *N*-(C₁₋₃-alkyl)-*N*-(C₃₋₆-cycloalkyl-C₁₋₃-alkyl-carbonyl)-amino, *N*-(C₁₋₃-alkyl)-*N*-(arylcarbonyl)-amino, *N*-(C₁₋₃-alkyl)-*N*-(aryl-C₁₋₃-alkyl-carbonyl)-amino, *N*-(C₁₋₃-alkyl)-*N*-(C₁₋₃-alkyloxy-carbonyl)-amino, *N*-(aminocarbonyl)-*N*-(C₁₋₃-alkyl)-amino, *N*-(C₁₋₃-alkyl-aminocarbonyl)-*N*-(C₁₋₃-alkyl)-amino, *N*-[di-(C₁₋₃-alkyl)aminocarbonyl]-*N*-(C₁₋₃-alkyl)-amino, *N*-(C₁₋₃-alkyl)-*N*-(C₁₋₃-alkyl-sulphonyl)-amino, *N*-(C₁₋₃-alkyl)-*N*-(arylsulphonyl)-amino or *N*-(C₁₋₃-alkyl)-*N*-(aryl-C₁₋₃-alkyl-sulphonyl)-amino group,

a 2-oxo-imidazolidin-1-yl, 2,4-dioxo-imidazolidin-1-yl, 2,5-dioxo-imidazolidin-1-yl or 2-oxo-hexahydropyrimidin-1-yl group wherein the nitrogen atom in the 3 position may be substituted in each case by a methyl or ethyl group,

a cyano, carboxy, C₁₋₃-alkyloxy-carbonyl, aminocarbonyl, C₁₋₃-alkyl-aminocarbonyl, di-(C₁₋₃-alkyl)-aminocarbonyl, pyrrolidin-1-yl-carbonyl, piperidin-1-yl-carbonyl, morpholin-4-yl-carbonyl, piperazin-1-yl-carbonyl or 4-(C₁₋₃-alkyl)-piperazin-1-yl-carbonyl group,

a C₁₋₃-alkyl-carbonyl or an arylcarbonyl group,

a carboxy-C₁₋₃-alkyl, C₁₋₃-alkyloxy-carbonyl-C₁₋₃-alkyl, cyano-C₁₋₃-alkyl, aminocarbonyl-C₁₋₃-alkyl, C₁₋₃-alkyl-aminocarbonyl-C₁₋₃-alkyl, di-(C₁₋₃-alkyl)-aminocarbonyl-C₁₋₃-alkyl, pyrrolidin-1-yl-carbonyl-C₁₋₃-alkyl,

piperidin-1-yl-carbonyl-C₁₋₃-alkyl, morpholin-4-yl-carbonyl-C₁₋₃-alkyl, piperazin-1-yl-carbonyl-C₁₋₃-alkyl or 4-(C₁₋₃-alkyl)-piperazin-1-yl-carbonyl-C₁₋₃-alkyl group,

a carboxy-C₁₋₃-alkyloxy, C₁₋₃-alkyloxy-carbonyl-C₁₋₃-alkyloxy, cyano-C₁₋₃-alkyloxy, aminocarbonyl-C₁₋₃-alkyloxy, C₁₋₃-alkyl-aminocarbonyl-C₁₋₃-alkyloxy, di-(C₁₋₃-alkyl)-aminocarbonyl-C₁₋₃-alkyloxy, pyrrolidin-1-yl-carbonyl-C₁₋₃-alkyl-oxy, piperidin-1-yl-carbonyl-C₁₋₃-alkyloxy, morpholin-4-yl-carbonyl-C₁₋₃-alkyl-oxy, piperazin-1-yl-carbonyl-C₁₋₃-alkyloxy or 4-(C₁₋₃-alkyl)-piperazin-1-yl-carbonyl-C₁₋₃-alkyloxy group,

a hydroxy-C₁₋₃-alkyl, C₁₋₃-alkyloxy-C₁₋₃-alkyl, amino-C₁₋₃-alkyl, C₁₋₃-alkyl-amino-C₁₋₃-alkyl, di-(C₁₋₃-alkyl)-amino-C₁₋₃-alkyl, pyrrolidin-1-yl-C₁₋₃-alkyl, piperidin-1-yl-C₁₋₃-alkyl, morpholin-4-yl-C₁₋₃-alkyl, piperazin-1-yl-C₁₋₃-alkyl, 4-(C₁₋₃-alkyl)-piperazin-1-yl-C₁₋₃-alkyl group,

a hydroxy-C₁₋₃-alkyloxy, C₁₋₃-alkyloxy-C₁₋₃-alkyloxy, C₁₋₃-alkylsulphanyl-C₁₋₃-alkyloxy, C₁₋₃-alkylsulphinyl-C₁₋₃-alkyloxy, C₁₋₃-alkylsulphonyl-C₁₋₃-alkyloxy, amino-C₁₋₃-alkyloxy, C₁₋₃-alkylamino-C₁₋₃-alkyloxy, di-(C₁₋₃-alkyl)-amino-C₁₋₃-alkyloxy, pyrrolidin-1-yl-C₁₋₃-alkyloxy, piperidin-1-yl-C₁₋₃-alkyloxy, morpholin-4-yl-C₁₋₃-alkyloxy, piperazin-1-yl-C₁₋₃-alkyloxy, 4-(C₁₋₃-alkyl)-piperazin-1-yl-C₁₋₃-alkyloxy group,

a mercapto, C₁₋₃-alkylsulphanyl, C₁₋₃-alkylsulphinyl, C₁₋₃-alkylsulphonyl, C₁₋₃-alkylsulphonyloxy, arylsulphonyloxy, trifluoromethylsulphanyl, trifluoromethylsulphinyl or trifluoromethylsulphonyl group,

a sulpho, aminosulphonyl, C₁₋₃-alkyl-aminosulphonyl, di-(C₁₋₃-alkyl)-aminosulphonyl, pyrrolidin-1-yl-sulphonyl, piperidin-1-yl-sulphonyl, morpholin-4-yl-sulphonyl, piperazin-1-yl-sulphonyl or 4-(C₁₋₃-alkyl)-piperazin-1-yl-sulphonyl group,

a methyl or methoxy group substituted by 1 to 3 fluorine atoms,

an ethyl or ethyloxy group substituted by 1 to 5 fluorine atoms,

a C₂₋₄-alkenyl or C₂₋₄-alkynyl group,

a C₃₋₄-alkenyloxy or C₃₋₄-alkynyloxy group,

a C₃₋₆-cycloalkyl or C₃₋₆-cycloalkyloxy group,

a C₃₋₆-cycloalkyl-C₁₋₃-alkyl or C₃₋₆-cycloalkyl-C₁₋₃-alkyloxy group or

an aryl, aryloxy, aryl-C₁₋₃-alkyl or aryl-C₁₋₃-alkyloxy group,

R¹¹ and R¹², which may be identical or different, in each case represent a hydrogen atom, a fluorine, chlorine, bromine or iodine atom, a C₁₋₃-alkyl, trifluoromethyl, hydroxy, C₁₋₃-alkyloxy or cyano group, or

R¹¹ together with R¹², if they are bound to adjacent carbon atoms, also represent a methylenedioxy, difluoromethylenedioxy or a straight-chain C₃₋₅-alkylene group and

R¹³ and R¹⁴, which may be identical or different, in each case represent a hydrogen atom, a fluorine, chlorine or bromine atom, a trifluoromethyl, C₁₋₃-alkyl or C₁₋₃-alkyloxy group,

a phenyl-C₁₋₄-alkyl group wherein the alkyl moiety is substituted by a cyano, carboxy, C₁₋₃-alkyloxy-carbonyl, aminocarbonyl, C₁₋₃-alkyl-aminocarbonyl, di-(C₁₋₃-alkyl)-aminocarbonyl, pyrrolidin-1-yl-carbonyl, piperidin-1-yl-carbonyl, morpholin-4-yl-carbonyl-group and the phenyl moiety is substituted by the groups R¹⁰ to R¹⁴, while R¹⁰ to R¹⁴ are as hereinbefore defined,

a phenyl group substituted by the groups R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴ are as hereinbefore defined,

a phenyl- C_{2-3} -alkenyl group wherein the phenyl moiety is substituted by the groups R^{10} to R^{14} , where R^{10} to R^{14} are as hereinbefore defined,

a phenyl- $(CH_2)_m$ -A- $(CH_2)_n$ -group wherein the phenyl moiety is substituted by R^{10} to R^{14} , where R^{10} to R^{14} are as hereinbefore defined and

A represents a carbonyl group, m represents the number 0, 1 or 2 and n represents the number 1, 2 or 3,

a phenylcarbonylmethyl group wherein the phenyl moiety is substituted by R^{10} to R^{14} , where R^{10} to R^{14} are as hereinbefore defined and the methyl moiety is substituted by a C_{1-3} -alkyl group,

a phenyl- $(CH_2)_m$ -B- $(CH_2)_n$ -group wherein the phenyl moiety is substituted by R^{10} to R^{14} , where R^{10} to R^{14} , m and n are as hereinbefore defined and

B denotes a methylene group which is substituted by a hydroxy, C_{1-3} -alkyloxy, amino, C_{1-3} -alkylamino, di- $(C_{1-3}$ -alkyl)-amino, mercapto, C_{1-3} -alkylsulphanyl, C_{1-3} -alkylsulphinyl or C_{1-3} -alkylsulphonyl group and is optionally additionally substituted by a methyl or ethyl group,

a naphthyl- C_{1-3} -alkyl group wherein the naphthyl moiety is substituted by the groups R^{10} to R^{14} , where R^{10} to R^{14} are as hereinbefore defined,

a naphthyl- $(CH_2)_m$ -A- $(CH_2)_n$ -group wherein the naphthyl moiety is substituted by R^{10} to R^{14} , where R^{10} to R^{14} , A, m and n are as hereinbefore defined,

a naphthyl- $(CH_2)_m$ -B- $(CH_2)_n$ -group wherein the naphthyl moiety is substituted by R^{10} to R^{14} , where R^{10} to R^{14} , B, m and n are as hereinbefore defined,

a [1,4]naphthoquinon-2-yl, chromen-4-on-3-yl, 1-oxoindan-2-yl, 1,3-dioxoindan-2-yl or 2,3-dihydro-3-oxo-benzofuran-2-yl group,

a heteroaryl-(CH₂)_m-A-(CH₂)_n group where A, m and n are as hereinbefore defined,

a heteroaryl-(CH₂)_m-B-(CH₂)_n group where B, m and n are as hereinbefore defined,

a C₁₋₆-alkyl-A-(CH₂)_n group where A and n are as hereinbefore defined,

a C₃₋₇-cycloalkyl-(CH₂)_m-A-(CH₂)_n group where A, m and n are as hereinbefore defined,

a C₃₋₇-cycloalkyl-(CH₂)_m-B-(CH₂)_n group where B, m and n are as hereinbefore defined,

an R²¹-A-(CH₂)_n-group wherein R²¹ denotes a C₁₋₃-alkyloxycarbonyl, aminocarbonyl, C₁₋₃-alkylaminocarbonyl, di-(C₁₋₃-alkyl)aminocarbonyl, pyrrolidin-1-yl-carbonyl, piperidin-1-yl-carbonyl or morpholin-4-yl-carbonyl, piperazin-1-yl-carbonyl, 4-methylpiperazin-1-yl-carbonyl or 4-ethylpiperazin-1-yl-carbonyl group and A and n are as hereinbefore defined,

a phenyl-(CH₂)_m-D-C₁₋₃-alkyl group wherein the phenyl moiety is substituted by the groups R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴ and m are as mentioned hereinbefore and D denotes an oxygen or sulphur atom, an imino, C₁₋₃-alkylimino, sulphinyl or sulphonyl group,

a naphthyl-(CH₂)_m-D-C₁₋₃-alkyl group wherein the naphthyl moiety is substituted by the groups R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴, D and m are as mentioned hereinbefore,

a C₂₋₆-alkyl group substituted by a group R_b, where

R_b is isolated from the cyclic nitrogen atom in the 1 position of the purine skeleton by at least two carbon atoms and

R_b denotes a hydroxy, C_{1-3} -alkyloxy, mercapto, C_{1-3} -alkylsulphanyl, C_{1-3} -alkylsulphinyl, C_{1-3} -alkylsulphonyl, amino, C_{1-3} -alkyl-carbonylamino, C_{3-6} -cycloalkyl-carbonyl-amino, arylcarbonylamino, C_{1-3} -alkylamino, di- $(C_{1-3}$ -alkyl)-amino, pyrrolidin-1-yl, piperidin-1-yl, morpholin-4-yl, piperazin-1-yl or 4- $(C_{1-3}$ -alkyl)-piperazin-1-yl group,

a C_{3-6} -cycloalkyl group,

or an amino or arylcarbonylamino group,

R^2 denotes a hydrogen atom,

a C_{1-8} -alkyl group,

a C_{3-8} -alkenyl group,

a C_{3-4} -alkenyl group which is substituted by a C_{1-2} -alkyloxy-carbonyl, aminocarbonyl, C_{1-3} -alkylamino-carbonyl, di- $(C_{1-3}$ -alkyl)-amino-carbonyl, pyrrolidin-1-ylcarbonyl, piperidin-1-ylcarbonyl or morpholin-4-ylcarbonyl group,

a C_{3-8} -alkynyl group,

a C_{3-6} -cycloalkyl group,

a C_{1-6} -alkyl group substituted by a group R_a , where R_a is as hereinbefore defined,

a phenyl group which is substituted by R^{10} to R^{14} , where R^{10} to R^{14} are as hereinbefore defined,

a C_{1-6} -alkyl group substituted by a phenyl group, wherein the phenyl ring is substituted by the groups R^{10} to R^{14} and R^{10} to R^{14} are as hereinbefore defined,

a phenyl-C₁₋₄-alkyl group wherein the alkyl moiety is substituted by a cyano, carboxy, C₁₋₃-alkyloxy-carbonyl, aminocarbonyl, C₁₋₃-alkyl-aminocarbonyl, di-(C₁₋₃-alkyl)-aminocarbonyl, pyrrolidin-1-yl-carbonyl, piperidin-1-yl-carbonyl, morpholin-4-yl-carbonyl group and the phenyl moiety is substituted by the groups R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴ are as hereinbefore defined,

a phenyl-C₂₋₃-alkenyl group wherein the phenyl moiety is substituted by R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴ are as hereinbefore defined,

a heteroaryl group,

a phenyl-(CH₂)_m-A or phenyl-(CH₂)_m-A-(CH₂)_n group wherein the phenyl moiety is substituted in each case by R¹⁰ to R¹⁴, while A, R¹⁰ to R¹⁴, m and n are as hereinbefore defined,

a phenylcarbonylmethyl group wherein the phenyl moiety is substituted by R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴ are as hereinbefore defined and the methyl moiety is substituted by a C₁₋₃-alkyl group,

a phenyl-(CH₂)_m-B or phenyl-(CH₂)_m-B-(CH₂)_n group wherein the phenyl moiety is substituted in each case by R¹⁰ to R¹⁴, while B, R¹⁰ to R¹⁴, m and n are as hereinbefore defined,

a naphthyl-C₁₋₃-alkyl group wherein the naphthyl moiety is substituted by the groups R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴ are as hereinbefore defined,

a naphthyl-(CH₂)_m-A or naphthyl-(CH₂)_m-A-(CH₂)_n group wherein the naphthyl moiety is substituted in each case by R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴, A, m and n are as hereinbefore defined,

a naphthyl-(CH₂)_m-B or naphthyl-(CH₂)_m-B-(CH₂)_n group wherein the naphthyl moiety is substituted in each case by R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴, B, m and n are as hereinbefore defined,

a heteroaryl-(CH₂)_m-A or heteroaryl-(CH₂)_m-A-(CH₂)_n group where A, m and n are as hereinbefore defined,

a heteroaryl-(CH₂)_m-B or heteroaryl-(CH₂)_m-B-(CH₂)_n group where B, m and n are as hereinbefore defined,

a C₁₋₆-alkyl-A or C₁₋₆-alkyl-A-(CH₂)_n group where A and n are as hereinbefore defined,

a C₃₋₇-cycloalkyl-(CH₂)_m-A or C₃₋₇-cycloalkyl-(CH₂)_m-A-(CH₂)_n group where A, m and n are as hereinbefore defined,

a C₃₋₇-cycloalkyl-(CH₂)_m-B or C₃₋₇-cycloalkyl-(CH₂)_m-B-(CH₂)_n group where B, m and n are as hereinbefore defined,

an R²¹-A-(CH₂)_n group wherein R²¹, A and n are as hereinbefore defined,

a phenyl-(CH₂)_m-D-C₁₋₃-alkyl group wherein the phenyl moiety is substituted by the groups R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴, D and m are as mentioned hereinbefore,

a naphthyl-(CH₂)_m-D-C₁₋₃-alkyl group wherein the naphthyl moiety is substituted by the groups R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴, D and m are as mentioned hereinbefore,

a C₁₋₆-alkyl group substituted by a group R_b, where R_b is as hereinbefore defined,

a cyano, carboxy, C₁₋₃-alkyloxy-carbonyl, aminocarbonyl, C₁₋₃-alkylamino-carbonyl, di-(C₁₋₃-alkyl)-amino-carbonyl, pyrrolidin-1-ylcarbonyl, piperidin-1-ylcarbonyl, morpholin-4-ylcarbonyl, piperazin-1-ylcarbonyl, 4-methylpiperazin-1-ylcarbonyl or 4-ethylpiperazin-1-ylcarbonyl group,

an amino, C₁₋₆-alkylamino or di-(C₁₋₆-alkyl)-amino group,

an amino group substituted by the groups R¹⁵ and R¹⁶ wherein

R¹⁵ denotes a hydrogen atom or a C₁₋₆-alkyl group and

R¹⁶ denotes a C₁₋₆-alkyl group which is substituted by R_a, where R_a is as hereinbefore defined,

an amino group substituted by the groups R¹⁵ and R¹⁷ wherein

R¹⁵ is as hereinbefore defined and

R¹⁷ denotes a C₂₋₆-alkyl group which is substituted by a hydroxy, C₁₋₃-alkyloxy, aryloxy, mercapto, C₁₋₃-alkylsulphanyl, C₁₋₃-alkylsulphinyl, C₁₋₃-alkylsulphonyl, C₁₋₃-alkylsulphonylamino, arylsulphanyl, arylsulphinyl, arylsulphonyl, arylsulphonylamino, C₁₋₃-alkyl-carbonylamino, C₃₋₆-cycloalkyl-carbonylamino, arylcarbonylamino, C₁₋₃-alkyl-oxycarbonylamino, aminocarbonylamino, C₁₋₃-alkyl-amino-carbonylamino, di-(C₁₋₃-alkyl)-aminocarbonylamino, amino, C₁₋₃-alkyl-amino, di-(C₁₋₃-alkyl)-amino, pyrrolidin-1-yl, piperidin-1-yl, morpholin-4-yl, piperazin-1-yl or 4-(C₁₋₃-alkyl)-piperazin-1-yl group,

a C₃₋₆-cycloalkylamino or *N*-(C₃₋₆-cycloalkyl)-*N*-(C₁₋₃-alkyl)-amino group,

a phenylamino or *N*-(phenyl)-*N*-(C₁₋₃-alkyl)-amino group wherein the phenyl moiety is substituted in each case by R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴ are as hereinbefore defined,

a phenyl-C₁₋₆-alkylamino or *N*-(phenyl-C₁₋₆-alkyl)-*N*-(C₁₋₃-alkyl)-amino group wherein the phenyl moiety is substituted in each case by R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴ are as hereinbefore defined,

a naphthylamino or *N*-(naphthyl)-*N*-(C₁₋₃-alkyl)-amino group,

a naphthyl-C₁₋₆-alkylamino or *N*-(naphthyl-C₁₋₆-alkyl)-*N*-(C₁₋₃-alkyl)-amino group,

a heteroaryl-amino or *N*-(heteroaryl)-*N*-(C₁₋₃-alkyl)-amino group,

a pyrrolidin-1-yl, piperidin-1-yl, homopiperidin-1-yl, morpholin-4-yl, homomorpholin-4-yl, piperazin-1-yl, 4-(C₁₋₃-alkyl)-piperazin-1-yl, homopiperazin-1-yl or 4-(C₁₋₃-alkyl)-homopiperazin-1-yl group, or

a C₁₋₆-alkyloxy, C₃₋₆-cycloalkyloxy or C₃₋₆-cycloalkyl-C₁₋₆-alkyloxy group,

a C₁₋₆-alkylsulphanyl, C₃₋₆-cycloalkylsulphanyl or C₃₋₆-cycloalkyl-C₁₋₆-alkylsulphanyl group,

a phenyloxy or phenylsulphanyl group wherein the phenyl moiety is substituted in each case by R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴ are as hereinbefore defined,

a phenyl-C₁₋₆-alkyloxy or phenyl-C₁₋₆-alkylsulphanyl group wherein the phenyl moiety is substituted in each case by R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴ are as hereinbefore defined,

a naphthyloxy or a naphthylsulphanyl group wherein the naphthyl moiety is substituted in each case by R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴ are as hereinbefore defined,

a naphthyl-C₁₋₆-alkyloxy or naphthyl-C₁₋₆-alkylsulphanyl group wherein the naphthyl moiety is substituted in each case by R¹⁰ to R¹⁴, where R¹⁰ to R¹⁴ are as hereinbefore defined,

a heteroaryloxy or heteroarylulphanyl group or

a heteroaryl-C₁₋₆-alkyloxy or heteroaryl-C₁₋₆-alkylsulphanyl group,

R^3 denotes a C_{1-8} -alkyl group,

a C_{1-4} -alkyl group substituted by the group R_c , where

R_c denotes a C_{3-7} -cycloalkyl group optionally substituted by one or two C_{1-3} -alkyl groups,

a C_{5-7} -cycloalkenyl group optionally substituted by one or two C_{1-3} -alkyl groups,

an aryl group or

a furanyl, thienyl, oxazolyl, isoxazolyl, thiazolyl, isothiazolyl, pyridyl, pyridazinyl, pyrimidyl or pyrazinyl group, while the abovementioned heterocyclic groups may each be substituted by one or two C_{1-3} -alkyl groups or by a fluorine, chlorine, bromine or iodine atom or by a trifluoromethyl, cyano or C_{1-3} -alkyloxy group,

a C_{3-8} -alkenyl group,

a C_{3-6} -alkenyl group substituted by a fluorine, chlorine or bromine atom or a trifluoromethyl group,

a C_{3-8} -alkynyl group,

an aryl group or

an aryl- C_{2-4} -alkenyl group,

and

R⁴ denotes an azetidin-1-yl or pyrrolidin-1-yl group which is substituted in the 3 position by an amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group and may additionally be substituted by one or two C₁₋₃-alkyl groups,

a piperidin-1-yl or hexahydroazepin-1-yl group which is substituted in the 3 position or in the 4 position by an amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)amino group and may additionally be substituted by one or two C₁₋₃-alkyl groups,

a 3-amino-piperidin-1-yl group wherein the piperidin-1-yl-moiety is additionally substituted by an aminocarbonyl, C₁₋₂-alkyl-aminocarbonyl, di-(C₁₋₂-alkyl)-aminocarbonyl, pyrrolidin-1-yl-carbonyl, (2-cyano-pyrrolidin-1-yl)-carbonyl, thiazolidin-3-yl-carbonyl, (4-cyano-thiazolidin-3-yl)carbonyl, piperidin-1-ylcarbonyl or morpholin-4-ylcarbonyl group,

a 3-amino-piperidin-1-yl group wherein the piperidin-1-yl moiety is additionally substituted in the 4 position or 5 position by a hydroxy or methoxy group,

a 3-amino-piperidin-1-yl group wherein the methylene group is replaced in the 2 position or 6 position by a carbonyl group,

a piperidin-1-yl or hexahydroazepin-1-yl- group substituted in the 3 position by an amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group, wherein two hydrogen atoms on the carbon skeleton of the piperidin-1-yl or hexahydroazepin-1-yl group are each replaced by a straight-chain alkylene bridge, this bridge containing 2 to 5 carbon atoms if the two hydrogen atoms are on the same carbon atom, or 1 to 4 carbon atoms if the hydrogen atoms are on adjacent carbon atoms, or 1 to 4 carbon atoms if the hydrogen atoms are on carbon atoms which are separated by one atom, or 1 to 3 carbon atoms if the hydrogen atoms are on carbon atoms which are separated by two atoms,

an azetidin-1-yl, pyrrolidin-1-yl, piperidin-1-yl or hexahydroazepin-1-yl group which is substituted by an amino- C_{1-3} -alkyl, C_{1-3} -alkylamino- C_{1-3} -alkyl or a di- $(C_{1-3}$ -alkyl)amino- C_{1-3} -alkyl group,

a piperazin-1-yl or [1,4]diazepan-1-yl group optionally substituted on the carbon skeleton by one or two C_{1-3} -alkyl groups,

a 3-imino-piperazin-1-yl, 3-imino-[1,4]diazepan-1-yl or 5-imino-[1,4]diazepan-1-yl group optionally substituted on the carbon skeleton by one or two C_{1-3} -alkyl groups,

a [1,4]diazepan-1-yl group optionally substituted by one or two C_{1-3} -alkyl groups, which is substituted in the 6 position by an amino group,

a C_{3-7} -cycloalkyl group which is substituted by an amino, C_{1-3} -alkylamino or di- $(C_{1-3}$ -alkyl)-amino group,

a C_{3-7} -cycloalkyl group which is substituted by an amino- C_{1-3} -alkyl, C_{1-3} -alkylamino- C_{1-3} -alkyl or a di- $(C_{1-3}$ -alkyl)amino- C_{1-3} -alkyl group,

a C_{3-7} -cycloalkyl- C_{1-2} -alkyl group wherein the cycloalkyl moiety is substituted by an amino, C_{1-3} -alkylamino or di- $(C_{1-3}$ -alkyl)-amino group,

a C_{3-7} -cycloalkyl- C_{1-2} -alkyl group wherein the cycloalkyl moiety is substituted by an amino- C_{1-3} -alkyl, C_{1-3} -alkylamino- C_{1-3} -alkyl or a di- $(C_{1-3}$ -alkyl)amino- C_{1-3} -alkyl group,

a C_{3-7} -cycloalkylamino group wherein the cycloalkyl moiety is substituted by an amino, C_{1-3} -alkylamino or di- $(C_{1-3}$ -alkyl)-amino group, while the two nitrogen atoms on the cycloalkyl moiety are separated from one another by at least two carbon atoms,

an N- $(C_{3-7}$ -cycloalkyl)-N- $(C_{1-3}$ -alkyl)-amino group wherein the cycloalkyl moiety is substituted by an amino, C_{1-3} -alkylamino or di- $(C_{1-3}$ -alkyl)-amino group,

while the two nitrogen atoms on the cycloalkyl moiety are separated from one another by at least two carbon atoms,

a C₃₋₇-cycloalkylamino group wherein the cycloalkyl moiety is substituted by an amino-C₁₋₃-alkyl, C₁₋₃-alkylamino-C₁₋₃-alkyl or a di-(C₁₋₃-alkyl)amino-C₁₋₃-alkyl group,

an N-(C₃₋₇-cycloalkyl)-N-(C₁₋₃-alkyl)-amino group wherein the cycloalkyl moiety is substituted by an amino-C₁₋₃-alkyl, C₁₋₃-alkylamino-C₁₋₃-alkyl or a di-(C₁₋₃-alkyl)amino-C₁₋₃-alkyl group,

a C₃₋₇-cycloalkyl-C₁₋₂-alkyl-amino group wherein the cycloalkyl moiety is substituted by an amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group,

an N-(C₃₋₇-cycloalkyl-C₁₋₂-alkyl)-N-(C₁₋₂-alkyl)-amino group wherein the cycloalkyl moiety is substituted by an amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group,

a C₃₋₇-cycloalkyl-C₁₋₂-alkyl-amino group wherein the cycloalkyl moiety is substituted by an amino-C₁₋₃-alkyl, C₁₋₃-alkylamino-C₁₋₃-alkyl or a di-(C₁₋₃-alkyl)amino-C₁₋₃-alkyl group,

an N-(C₃₋₇-cycloalkyl-C₁₋₂-alkyl)-N-(C₁₋₂-alkyl)-amino group wherein the cycloalkyl moiety is substituted by an amino-C₁₋₃-alkyl, C₁₋₃-alkylamino-C₁₋₃-alkyl or a di-(C₁₋₃-alkyl)amino-C₁₋₃-alkyl group,

an R¹⁹-C₂₋₄-alkylamino group wherein R¹⁹ is separated from the nitrogen atom of the C₂₋₄-alkylamino moiety by at least two carbon atoms and

R¹⁹ denotes an amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)-amino group,

an R¹⁹-C₂₋₄-alkylamino group wherein the nitrogen atom of the C₂₋₄-alkylamino moiety is substituted by a C₁₋₃-alkyl group and R¹⁹ is separated from the

nitrogen atom of the C₂₋₄-alkylamino moiety by at least two carbon atoms, where R¹⁹ is as hereinbefore defined,

an amino group substituted by the group R²⁰ wherein

R²⁰ denotes an azetidin-3-yl, azetidin-2-ylmethyl, azetidin-3-ylmethyl, pyrrolidin-3-yl, pyrrolidin-2-ylmethyl, pyrrolidin-3-ylmethyl, piperidin-3-yl, piperidin-4-yl, piperidin-2-ylmethyl, piperidin-3-ylmethyl or piperidin-4-ylmethyl group, while the groups mentioned for R²⁰ may each be substituted by one or two C₁₋₃-alkyl groups,

an amino group substituted by the group R²⁰ and a C₁₋₃-alkyl group wherein R²⁰ is as hereinbefore defined, while the groups mentioned for R²⁰ may each be substituted by one or two C₁₋₃-alkyl groups,

an R¹⁹-C₃₋₄-alkyl group wherein the C₃₋₄-alkyl moiety is straight-chained and may additionally be substituted by one or two C₁₋₃-alkyl groups, where R¹⁹ is as hereinbefore defined,

a 3-amino-2-oxo-piperidin-5-yl or 3-amino-2-oxo-1-methyl-piperidin-5-yl group,

a pyrrolidin-3-yl, piperidin-3-yl, piperidin-4-yl, hexahydroazepin-3-yl or hexahydroazepin-4-yl group which is substituted in the 1 position by an amino, C₁₋₃-alkylamino or di-(C₁₋₃-alkyl)amino group,

or an azetidin-2-yl-C₁₋₂-alkyl, azetidin-3-yl-C₁₋₂-alkyl, pyrrolidin-2-yl-C₁₋₂-alkyl, pyrrolidin-3-yl, pyrrolidin-3-yl-C₁₋₂-alkyl, piperidin-2-yl-C₁₋₂-alkyl, piperidin-3-yl, piperidin-3-yl-C₁₋₂-alkyl, piperidin-4-yl or piperidin-4-yl-C₁₋₂-alkyl group, while the abovementioned groups may each be substituted by one or two C₁₋₃-alkyl groups,

while by the aryl groups mentioned in the definition of the above groups are meant phenyl or naphthyl groups, which may be mono- or disubstituted by R_n independently of one another, where the substituents are identical or different

and R_h denotes a fluorine, chlorine, bromine or iodine atom, a trifluoromethyl, cyano, nitro, amino, aminocarbonyl, aminosulphonyl, methylsulphonyl, acetylamino, methylsulphonylamino, C_{1-3} -alkyl, cyclopropyl, ethenyl, ethynyl, hydroxy, C_{1-3} -alkyloxy, difluoromethoxy or trifluoromethoxy group,

by the heteroaryl groups mentioned in the definitions of the abovementioned groups are meant a pyrrolyl, furanyl, thienyl, pyridyl, indolyl, benzofuranyl, benzothiophenyl, quinolinyl or isoquinolinyl group,

or a pyrrolyl, furanyl, thienyl or pyridyl group wherein one or two methyne groups are replaced by nitrogen atoms,

or an indolyl, benzofuranyl, benzothiophenyl, quinolinyl or isoquinolinyl group wherein one to three methyne groups are replaced by nitrogen atoms,

or a 1,2-dihydro-2-oxo-pyridinyl, 1,4-dihydro-4-oxo-pyridinyl, 2,3-dihydro-3-oxo-pyridazinyl, 1,2,3,6-tetrahydro-3,6-dioxo-pyridazinyl, 1,2-dihydro-2-oxo-pyrimidinyl, 3,4-dihydro-4-oxo-pyrimidinyl, 1,2,3,4-tetrahydro-2,4-dioxo-pyrimidinyl, 1,2-dihydro-2-oxo-pyrazinyl, 1,2,3,4-tetrahydro-2,3-dioxo-pyrazinyl, 2,3-dihydro-2-oxo-indolyl, 2,3-dihydrobenzofuranyl, 2,3-dihydro-2-oxo-1*H*-benzimidazolyl, 2,3-dihydro-2-oxo-benzoxazolyl, 1,2-dihydro-2-oxo-quinolinyl, 1,4-dihydro-4-oxo-quinolinyl, 1,2-dihydro-1-oxo-isoquinolinyl, 1,4-dihydro-4-oxo-cinnolinyl, 1,2-dihydro-2-oxo-quinazolinyl, 3,4-dihydro-4-oxo-quinazolinyl, 1,2,3,4-tetrahydro-2,4-dioxo-quinazolinyl, 1,2-dihydro-2-oxoquinoxaliny, 1,2,3,4-tetrahydro-2,3-dioxo-quinoxaliny, 1,2-dihydro-1-oxo-phthalazinyl, 1,2,3,4-tetrahydro-1,4-dioxo-phthalazinyl, chromanyl, cumariny, 2,3-dihydro-benzo[1,4]dioxiny or 3,4-dihydro-3-oxo-2*H*-benzo[1,4]oxaziny group,

while the abovementioned heteroaryl groups may be substituted by R^{10} to R^{14} , where R^{10} to R^{14} are as hereinbefore defined,

and, unless otherwise stated, the abovementioned alkyl, alkenyl and alkynyl groups may be straight-chain or branched,

as well as the derivatives which are N-oxidised or methylated or ethylated at the cyclic nitrogen atom in the 3 position or 9 position of the hypoxanthine skeleton,

as well as the derivatives wherein the 6-oxo group of the hypoxanthine skeleton is replaced by a thioxo group,

with the proviso that the compounds

8-(piperidin-4-ylmethyl)-7-(4-fluorobenzyl)-1,7-dihydro-purin-6-one and

8-(1-methyl-piperidin-4-ylmethyl)-7-(4-fluorobenzyl)-1,7-dihydro-purin-6-one

are excluded,

the tautomers, enantiomers, diastereomers, the mixtures thereof, the prodrugs thereof and the salts thereof.

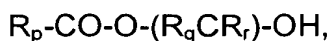
Compounds which contain a group that can be cleaved *in vivo* are prodrugs of the corresponding compounds wherein this group that can be cleaved *in vivo* has been cleaved.

The carboxy groups mentioned in the definition of the abovementioned groups may be replaced by a group which can be converted into a carboxy group *in vivo* or by a group which is negatively charged under physiological conditions,

and furthermore the amino and imino groups mentioned in the definition of the abovementioned groups may be substituted by a group which can be cleaved *in vivo*. Such groups are described for example in WO 98/46576 and by N.M. Nielsen *et al.* in International Journal of Pharmaceutics 39, 75-85 (1987).

By a group which can be converted *in vivo* into a carboxy group is meant, for example, a hydroxymethyl group, a carboxy group esterified with an alcohol

wherein the alcohol moiety is preferably a C₁₋₆-alkanol, a phenyl-C₁₋₃-alkanol, a C₃₋₉-cycloalkanol, while a C₅₋₈-cycloalkanol may additionally be substituted by one or two C₁₋₃-alkyl groups, a C₅₋₈-cycloalkanol wherein a methylene group in the 3 or 4 position is replaced by an oxygen atom or by an imino group optionally substituted by a C₁₋₃-alkyl, phenyl-C₁₋₃-alkyl, phenyl-C₁₋₃-alkyloxycarbonyl or C₂₋₆-alkanoyl group and the cycloalkanol moiety may additionally be substituted by one or two C₁₋₃-alkyl groups, a C₄₋₇-cycloalkenol, a C₃₋₅-alkenol, a phenyl-C₃₋₅-alkenol, a C₃₋₅-alkynol or phenyl-C₃₋₅-alkynol with the proviso that no bonds to the oxygen atom start from a carbon atom which carries a double or triple bond, a C₃₋₈-cycloalkyl-C₁₋₃-alkanol, a bicycloalkanol with a total of 8 to 10 carbon atoms which may additionally be substituted in the bicycloalkyl moiety by one or two C₁₋₃-alkyl groups, a 1,3-dihydro-3-oxo-1-isobenzofuranol or an alcohol of formula



wherein

R_p denotes a C₁₋₈-alkyl, C₅₋₇-cycloalkyl, C₁₋₈-alkyloxy, C₅₋₇-cycloalkyloxy, phenyl or phenyl-C₁₋₃-alkyl group,

R_q denotes a hydrogen atom, a C₁₋₃-alkyl, C₅₋₇-cycloalkyl or phenyl group and

R_r denotes a hydrogen atom or a C₁₋₃-alkyl group,

by a group which is negatively charged under physiological conditions is meant, for example, a tetrazol-5-yl, phenylcarbonylaminocarbonyl, trifluoromethylcarbonylaminocarbonyl, C₁₋₆-alkylsulphonylamino, phenylsulphonylamino, benzylsulphonylamino, trifluoromethylsulphonylamino, C₁₋₆-alkylsulphonylaminocarbonyl, phenylsulphonylaminocarbonyl, benzylsulphonylaminocarbonyl or perfluoro-C₁₋₆-alkylsulphonylaminocarbonyl group

and by a group which can be cleaved *in vivo* from an imino or amino group is meant, for example, a hydroxy group, an acyl group such as a phenylcarbonyl group optionally mono- or disubstituted by fluorine, chlorine, bromine or iodine atoms, by C₁₋₃-alkyl or C₁₋₃-alkyloxy groups, while the substituents may be identical or different, a pyridinoyl group or a C₁₋₁₆-alkanoyl group such as the formyl, acetyl, propionyl, butanoyl, pentanoyl or hexanoyl group, a 3,3,3-trichloropropionyl or allyloxycarbonyl group, a C₁₋₁₆-alkyloxycarbonyl or C₁₋₁₆-alkylcarbonyloxy group, wherein hydrogen atoms may be wholly or partially replaced by fluorine or chlorine atoms such as the methoxycarbonyl, ethoxycarbonyl, propoxycarbonyl, isopropoxycarbonyl, butoxycarbonyl, tert.butoxycarbonyl, pentoxycarbonyl, hexoxycarbonyl, octyloxycarbonyl, nonyloxycarbonyl, decyloxycarbonyl, undecyloxycarbonyl, dodecyloxycarbonyl, hexadecyloxycarbonyl, methylcarbonyloxy, ethylcarbonyloxy, 2,2,2-trichloroethylcarbonyloxy, propylcarbonyloxy, isopropylcarbonyloxy, butylcarbonyloxy, tert.butylcarbonyloxy, pentylcarbonyloxy, hexylcarbonyloxy, octylcarbonyloxy, nonylcarbonyloxy, decylcarbonyloxy, undecylcarbonyloxy, dodecylcarbonyloxy or hexadecylcarbonyloxy group, a phenyl-C₁₋₆-alkyloxycarbonyl group such as the benzyloxycarbonyl, phenylethoxycarbonyl or phenylpropoxycarbonyl group, a 3-amino-propionyl group wherein the amino group may be mono- or disubstituted by C₁₋₆-alkyl or C₃₋₇-cycloalkyl groups and the substituents may be identical or different, a C₁₋₃-alkylsulphonyl-C₂₋₄-alkyloxycarbonyl, C₁₋₃-alkyloxy-C₂₋₄-alkyloxy-C₂₋₄-alkyloxycarbonyl, R_p-CO-O-(R_qCR_r)-O-CO-, C₁₋₆-alkyl-CO-NH-(R_sCR_t)-O-CO- or C₁₋₆-alkyl-CO-O-(R_sCR_t)-(R_sCR_t)-O-CO-group, wherein R_p to R_r are as hereinbefore defined,

R_s and R_t, which may be identical or different, denote hydrogen atoms or C₁₋₃-alkyl groups.

Moreover, the saturated alkyl and alkyloxy moieties which contain more than 2 carbon atoms mentioned in the foregoing definitions and those that follow, unless otherwise stated, also include the branched isomers thereof such as, for example, the isopropyl, tert.butyl, isobutyl group, etc.

One sub-group deserving special mention relates to those compounds of general formula I wherein R^1 , R^2 and R^3 are as hereinbefore defined and

R^4 denotes a pyrrolidin-1-yl group which is substituted in the 3 position by an amino group,

a piperidin-1-yl group which is substituted in the 3 position by an amino group,

a piperidin-3-yl or piperidin-4-yl group,

a hexahydroazepin-1-yl group which is substituted in the 3 position or 4 position by an amino group,

a piperazin-1-yl or [1,4]diazepan-1-yl group,

a (2-aminocyclohexyl)amino group,

a cyclohexyl group which is substituted in the 3 position by an amino group,

or an N-(2-aminoethyl)-methylamino or an N-(2-aminoethyl)-ethylamino group,

the tautomers, enantiomers, diastereomers, the mixtures thereof, the prodrugs and the salts thereof.

Preferred compounds of the above general formula I are those wherein

R^1 denotes a hydrogen atom,

a C_{1-6} -alkyl group,

a C_{3-6} -alkenyl group,

a C_{3-4} -alkynyl group,

a C₃₋₆-cycloalkylmethyl group,

a phenyl-C₁₋₃-alkyl group wherein the phenyl moiety is substituted by R¹⁰ and R¹¹, where

R¹⁰ denotes a hydrogen atom, a fluorine, chlorine or bromine atom,

a methyl or trifluoromethyl group,

a cyano, aminocarbonyl, dimethylaminocarbonyl or methylsulphonyl group,

an amino, acetylamino or methylsulphonylamino group,

a hydroxy, methoxy, difluoromethoxy, trifluoromethoxy, carboxymethoxy, methoxycarbonylmethoxy, ethyloxycarbonylmethoxy, aminocarbonylmethoxy, methylaminocarbonylmethoxy, ethylamino-carbonylmethoxy or dimethylaminocarbonylmethoxy group and

R¹¹ denotes a hydrogen atom, a fluorine or chlorine atom,

or a methyl or methoxy group,

a naphthylmethyl group,

a heteroarylmethyl group where the term

heteroaryl denotes a furanyl, thienyl, oxazolyl, isoxazolyl, thiazolyl, pyridyl, pyrimidinyl, pyrazinyl, quinolinyl, isoquinolinyl or quinazolinyl group and the abovementioned heteroaryl groups may be substituted by one or two methyl groups,

a phenylcarbonylmethyl group wherein the phenyl moiety is substituted by R^{10} and R^{11} , where R^{10} and R^{11} are as hereinbefore defined,

a furanylcabonylmethyl, thienylcarbonylmethyl or pyridylcarbonylmethyl group,

or a 2-oxo-propyl or cyclohexylcarbonylmethyl group,

R^2 denotes a hydrogen atom,

a C_{1-6} -alkyl group,

a C_{3-6} -alkenyl group,

a C_{3-4} -alkynyl group,

a C_{3-6} -cycloalkyl or C_{3-6} -cycloalkyl- C_{1-3} -alkyl group,

a phenyl group which is substituted by R^{10} and R^{11} , where R^{10} and R^{11} are as hereinbefore defined,

a phenyl- C_{1-3} -alkyl group wherein the phenyl moiety is substituted by R^{10} and R^{11} , where R^{10} and R^{11} are as hereinbefore defined,

a phenyl- C_{2-3} -alkenyl group wherein the phenyl moiety is substituted by R^{10} and R^{11} , where R^{10} and R^{11} are as hereinbefore defined,

a furanyl, thienyl or pyridyl group,

a furanyl- C_{1-3} -alkyl, thienyl- C_{1-3} -alkyl or pyridyl- C_{1-3} -alkyl group,

an amino, C_{1-4} -alkylamino or di- $(C_{1-4}$ -alkyl)-amino group,

an amino group substituted by the groups R^{15} and R^{16} wherein

R^{15} denotes a hydrogen atom or a methyl or ethyl group and

R^{16} denotes a C_{1-4} -alkyl group which is substituted by a cyano, carboxy, methoxycarbonyl, ethyloxycarbonyl, aminocarbonyl, methylaminocarbonyl, dimethylaminocarbonyl, ethylaminocarbonyl, diethylaminocarbonyl, pyrrolidin-1-ylcarbonyl or morpholin-4-ylcarbonyl group,

an amino group substituted by the groups R^{15} and R^{17} wherein

R^{15} is as hereinbefore defined and

R^{17} denotes a straight-chain C_{2-4} -alkyl group which is terminally substituted in each case by an amino, methylamino, dimethylamino, acetylamino, ethyloxycarbonylamino, phenylcarbonylamino, methylsulphonylamino, phenylsulphonylamino, hydroxy, methoxy, phenoxy, methylsulphanyl or phenylsulphanyl group,

a pyrrolidin-1-yl, piperidin-1-yl, morpholin-4-yl, piperazin-1-yl or 4-methylpiperazin-1-yl group,

a C_{3-6} -cycloalkylamino or C_{3-6} -cycloalkyl- C_{1-3} -alkylamino group,

a phenylamino group,

a phenyl- C_{1-3} -alkylamino group wherein the phenyl moiety is substituted by R^{10} and R^{11} , where R^{10} and R^{11} are as hereinbefore defined,

a naphthylmethylamino group,

a heteroaryl- C_{1-2} -alkylamino group, where the term heteroaryl is as hereinbefore defined, or

a methylsulphanyl, benzylsulphanyl or (2-phenylethyl)sulphanyl group,

R^3 denotes a C_{4-6} -alkenyl group,

a C_{3-4} -alkenyl group which is substituted by a fluorine, chlorine or bromine atom or a trifluoromethyl group,

a 2-butyne-1-yl group or

a methyl group substituted by the group R_c , where

R_c denotes a 1-cyclopenten-1-yl, 1-cyclohexen-1-yl or cyclohexyl group,

a phenyl group optionally substituted by a fluorine, chlorine, bromine or iodine atom, by a methyl, trifluoromethyl, cyano, methoxy, difluoromethoxy or trifluoromethoxy group,

a phenyl group which is substituted by two fluorine atoms,

a naphthyl group or

a furanyl, thienyl, or pyridyl group,

and

R^4 denotes a pyrrolidin-1-yl group which is substituted in the 3 position by an amino group,

a piperidin-1-yl group which is substituted in the 3 position by an amino group,

a piperidin-3-yl or piperidin-4-yl group,

a hexahydroazepin-1-yl group which is substituted in the 3 position or 4 position by an amino group,

a piperazin-1-yl or [1,4]diazepan-1-yl group,

a (2-aminocyclohexyl)amino group,

a cyclohexyl group which is substituted in the 3 position by an amino group, or

an N-(2-aminoethyl)-methylamino or an N-(2-aminoethyl)-ethylamino group,

while unless otherwise stated, the abovementioned alkyl, alkenyl and alkynyl groups may be straight-chain or branched,

the tautomers, enantiomers, diastereomers, the mixtures thereof and the salts thereof.

Particularly preferred compounds of the above general formula I are those wherein

R^1 denotes a hydrogen atom,

a methyl, benzyl or 2-phenylethyl group or

a phenylcarbonylmethyl group,

R^2 denotes a hydrogen atom,

a methyl or 2-phenylethyl group,

a methylsulphanyl, benzylsulphanyl or (2-phenylethyl)sulphanyl group,

an amino, methylamino, dimethylamino, isopropylamino, cyclohexylamino- or (cyclohexylmethyl)amino group,

a benzylamino, fluorobenzylamino or (2-phenylethyl)amino group or

a piperidin-1-yl group,

R³ denotes a benzyl or 3-methyl-but-2-en-1-yl group

and

R⁴ denotes a (3-amino-piperidin-1-yl) group,

the tautomers, enantiomers, diastereomers, the mixtures thereof and the salts thereof,

but particularly the compounds

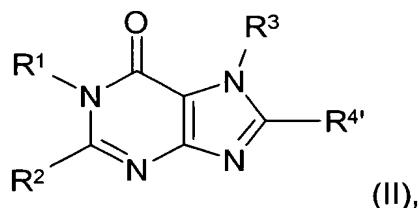
- (1) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-benzylamino-1-methyl-1,7-dihydro-purin-6-one,
- (2) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-(4-fluoro-benzylamino)-1-methyl-1,7-dihydro-purin-6-one,
- (3) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(2-phenylethyl)amino]-1,7-dihydro-purin-6-one,
- (4) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-isopropylamino-1-methyl-1,7-dihydro-purin-6-one,
- (5) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-methylamino-1,7-dihydro-purin-6-one,
- (6) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-cyclohexylamino-1-methyl-1,7-dihydro-purin-6-one,
- (7) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-[(cyclohexylmethyl)amino]-1-methyl-1,7-dihydro-purin-6-one,

- (8) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(piperidin-1-yl)-1,7-dihydro-purin-6-one,
- (9) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-dimethylamino-1-methyl-1,7-dihydro-purin-6-one,
- (10) 2-amino-8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-1,7-dihydro-purin-6-one,
- (11) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(3-methyl-but-2-en-1-yl)-1,7-dihydro-purin-6-one,
- (12) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-methyl-1,7-dihydro-purin-6-one,
- (13) 8-(3-amino-piperidin-1-yl)-1-methyl-7-(3-methyl-but-2-en-1-yl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one,
- (14) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-1,7-dihydro-purin-6-one,
- (15) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-methylsulphanyl-1,7-dihydro-purin-6-one,
- (16) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-benzylsulphanyl-1,7-dihydro-purin-6-one,
- (17) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-methylsulphanyl-1,7-dihydro-purin-6-one,
- (18) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-methylsulphanyl-1-(2-phenyl-2-oxo-ethyl)-1,7-dihydro-purin-6-one,
- (19) 8-(3-amino-piperidin-1-yl)-1-benzyl-7-benzyl-2-methylsulphanyl-1,7-dihydro-purin-6-one,

- (20) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-methylsulphanyl-1-(2-phenylethyl)-1,7-dihydro-purin-6-one and
- (21) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2-oxo-2-phenyl-ethyl)-1,7-dihydro-purin-6-one

as well as the tautomers, enantiomers, diastereomers, the mixtures thereof and the salts thereof.

The compounds of general formula I can be prepared by deprotecting a compound of general formula



wherein R^1 , R^2 and R^3 are as hereinbefore defined and $R^{4'}$ denotes one of the groups mentioned hereinbefore for R^4 which contain an imino, amino or alkylamino group, while the imino, amino or alkylamino group is substituted by a protective group, optionally followed by subsequent alkylation of the imino, amino or C_{1-3} -alkylamino group.

The liberating of an amino group from a protected precursor is a standard reaction in synthetic organic chemistry. There are many examples of suitable protective groups. A summary of the chemistry of protective groups can be found in Theodora W. Greene and Peter G.M. Wuts, *Protective Groups in Organic Synthesis*, Second Edition, 1991, published by John Wiley and Sons, and in Philip J. Kocienski, *Protecting Groups*, published by Georg Thieme, 1994.

The following are examples of protective groups:

the tert.-butoxycarbonyl group which can be cleaved by treating with an acid such as for example trifluoroacetic acid or hydrogen chloride in the presence of a solvent such as for example methylene chloride, ethyl acetate or dioxane at temperatures between 0°C and the boiling temperature of the solvent used,

the 2,2,2-trichloroethoxycarbonyl group which can be cleaved by treating with metals such as for example zinc or cadmium in a solvent such as acetic acid or a mixture of tetrahydrofuran and a weak aqueous acid at temperatures between 0°C and the boiling temperature of the solvent used and

the carbobenzyloxycarbonyl group which can be cleaved for example by hydrogenolysis in the presence of a noble metal catalyst such as for example palladium-charcoal and a solvent such as for example alcohols, ethyl acetate, dioxane, tetrahydrofuran or mixtures of these solvents at temperatures between 0°C and the boiling point of the solvent, by treating with boron tribromide in methylene chloride at temperatures between -20°C and ambient temperature, or by treating with aluminium chloride/anisole at temperatures between 0°C and ambient temperature.

The optional subsequent introduction of a C₁₋₃-alkyl group may be done by alkylation or reductive alkylation.

The subsequent alkylation is optionally carried out in a solvent or mixture of solvents such as methylene chloride, dimethylformamide, benzene, toluene, chlorobenzene, tetrahydrofuran, benzene/tetrahydrofuran or dioxane with an alkylating agent such as a corresponding halide or sulphonic acid ester, e.g. with methyl iodide, ethyl bromide, dimethyl sulphate, optionally in the presence of a tertiary organic base or in the presence of an inorganic base, conveniently at temperatures between 0 and 150°C, preferably at temperatures between 0 and 100°C.

The subsequent reductive alkylation is carried out with a corresponding carbonyl compound such as formaldehyde, acetaldehyde, propionaldehyde or

acetone in the presence of a complex metal hydride such as sodium borohydride, lithium borohydride, sodium triacetoxyborohydride or sodium cyanoborohydride, conveniently at a pH of 6-7 and at ambient temperature or in the presence of a hydrogenation catalyst, e.g. with hydrogen in the presence of palladium/charcoal, under a hydrogen pressure of 1 to 5 bar. The methylation may also be carried out in the presence of formic acid as reducing agent at elevated temperatures, e.g. at temperatures between 60 and 120°C.

The compounds of general formula I obtained may be resolved into their enantiomers and/or diastereomers. Thus, for example, cis/trans mixtures may be resolved into their cis and trans isomers, and compounds with at least one optically active carbon atom may be separated into their enantiomers.

Thus, for example, the cis/trans mixtures may be resolved by chromatography into the cis and trans isomers thereof, the compounds of general formula I obtained which occur as racemates may be separated by methods known per se (cf. Allinger N. L. and Eliel E. L. in "Topics in Stereochemistry", Vol. 6, Wiley Interscience, 1971) into their optical antipodes and compounds of general formula I with at least 2 asymmetric carbon atoms may be resolved into their diastereomers on the basis of their physical-chemical differences using methods known per se, e.g. by chromatography and/or fractional crystallisation, and, if these compounds are obtained in racemic form, they may subsequently be resolved into the enantiomers as mentioned above.

The enantiomers are preferably separated by column separation on chiral phases or by recrystallisation from an optically active solvent or by reacting with an optically active substance which forms salts or derivatives such as e.g. esters or amides with the racemic compound, particularly acids and the activated derivatives or alcohols thereof, and separating the diastereomeric mixture of salts or derivatives thus obtained, e.g. on the basis of their differences in solubility, whilst the free antipodes may be released from the pure diastereomeric salts or derivatives by the action of suitable agents. Optically active acids in common use are e.g. the D- and L-forms of tartaric acid or dibenzoyltartaric acid, di-o-tolyltartaric acid, malic acid, mandelic acid,

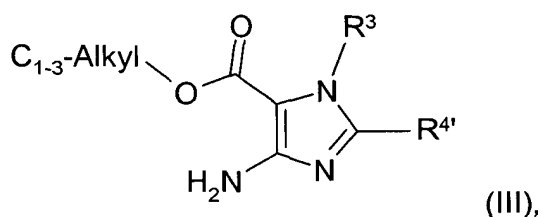
camphorsulphonic acid, glutamic acid, aspartic acid or quinic acid. An optically active alcohol may be for example (+) or (-)-menthol and an optically active acyl group in amides, for example, may be a (+)-or (-)-menthyloxycarbonyl.

Furthermore, the compounds of formula I obtained may be converted into the salts thereof, particularly for pharmaceutical use into the physiologically acceptable salts with inorganic or organic acids. Acids which may be used for this purpose include for example hydrochloric acid, hydrobromic acid, sulphuric acid, methane sulphonic acid, phosphoric acid, fumaric acid, succinic acid, lactic acid, citric acid, tartaric acid or maleic acid.

Moreover, if the new compounds of formula I thus obtained contain a carboxy group, they may subsequently, if desired, be converted into the salts thereof with inorganic or organic bases, particularly for pharmaceutical use into the physiologically acceptable salts thereof. Suitable bases for this purpose include for example sodium hydroxide, potassium hydroxide, arginine, cyclohexylamine, ethanolamine, diethanolamine and triethanolamine.

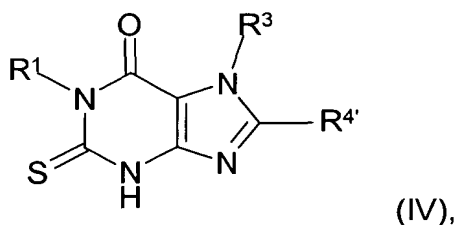
The starting compounds of general formula II may be prepared by generally known methods and according to processes described for example in Examples I to XIV.

Thus, compounds of general formula II wherein R^1 , R^3 and $R^{4'}$ are as hereinbefore defined and R^2 denotes a hydrogen atom, may be prepared for example by reacting a compound of formula



with a formimido-C₁₋₃-alkyl ester, optionally followed by alkylation at N-1 with a suitable alkylating agent.

Compounds of general formula II wherein R¹, R³ and R^{4'} are as hereinbefore defined and R² denotes an alkylsulphanyl, an alkenylsulphanyl or alkynylsulphanyl group may be obtained for example by reacting compounds of general formula III with a suitable isothiocyanate and subsequently cyclising them to form compounds of general formula



followed by alkylation of the sulphur atom.

If an acyl mustard oil is used, such as for example ethoxycarbonylisothiocyanate, first of all compounds of general formula IV, wherein R¹ denotes a hydrogen atom are obtained, which can be converted into the desired compounds by subsequent alkylation at the sulphur atom and at N-1.

Compounds of general formula II wherein R¹, R³ and R^{4'} are as hereinbefore defined and R² denotes an alkylsulphanyl group can be oxidised to form compounds of formula II' wherein R^{2'} denotes an alkylsulphinyl or an alkylsulphonyl group.

The abovementioned compounds of general formula II' are starting materials for preparing the following compounds of formula II.

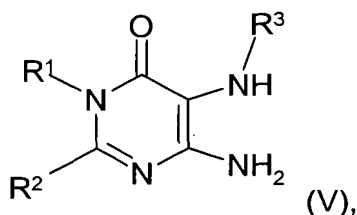
Reaction with alcohols and phenols yields compounds wherein the group R² is linked to the purine system via an oxygen atom,

Reaction with thiols and thiophenols leads to compounds wherein R^2 is linked to the purine system via a sulphur atom,

Reaction with amines leads to compounds wherein R^2 is bound to the purine system via a nitrogen atom and

Reaction with organometallic compounds such as for example Grignard reagents, alkyl- or aryl-lithium compounds or reaction with CH-acid compounds such as for example esters, nitriles or ketones leads to compounds wherein R^2 is linked to the purine system via a carbon atom.

Another method of obtaining compounds of general formula II consists, for example, in converting compounds of general formula



wherein R^1 , R^2 and R^3 are as hereinbefore defined, into compounds of general formula II.

For example, compounds of general formula V are converted into compounds of general formula II wherein the group $R^{4'}$ is bound to the purine system via a nitrogen atom, by reacting with an orthoformate, subsequently brominating the resulting purine at C-8 and then reacting with a corresponding amine.

For example, compounds of general formula V are converted into compounds of general formula II wherein the group $R^{4'}$ is bound to the C-8 atom of the purine via a C-atom by reaction with a reactive derivative of a carboxylic acid $R^{4'}-\text{COOH}$, where $R^{4'}$ is as hereinbefore defined, and subsequent cyclisation.

As already mentioned hereinbefore, the compounds of general formula I according to the invention and the physiologically acceptable salts thereof have valuable pharmacological properties, particularly an inhibiting effect on the enzyme DPP-IV.

The biological properties of the new compounds were investigated as follows:

The ability of the substances and their corresponding salts to inhibit the DPP-IV activity can be demonstrated in a test set-up in which an extract of human colon carcinoma cell line Caco-2 is used as the DPP IV source. The differentiation of the cells in order to induce the DPP-IV expression was carried out as described by Reiher et al. in an article entitled "Increased expression of intestinal cell line Caco-2", which appeared in Proc. Natl. Acad. Sci. Vol. 90, pages 5757-5761 (1993). The cell extract was obtained from cells solubilised in a buffer (10mM Tris HCl, 0.15 M NaCl, 0.04 t.i.u. aprotinin, 0.5% Nonidet-P40, pH 8.0) by centrifuging at 35,000 g for 30 minutes at 4°C (to remove cell debris).

The DPP-IV assay was carried out as follows:

50 µl substrate solution (AFC; AFC is amido-4-trifluoromethylcoumarin), final concentration 100 µM, were placed in black microtitre plates. 20 µl of assay buffer (final concentrations 50 mM Tris HCl pH 7.8, 50 mM NaCl, 1 % DMSO) was pipetted in. The reaction was started by adding 30 µl of solubilised Caco-2 protein (final concentration 0.14 µg of protein per well). The test substances to be investigated were typically added prediluted in 20 µl, and the volume of assay buffer was then reduced accordingly. The reaction was carried out at ambient temperature, incubating for 60 minutes. Then the fluorescence was measured in a Victor 1420 Multilabel Counter, the excitation wavelength being 405 nm and the emission wavelength being 535 nm. Blank readings (corresponding to 0 % activity) were obtained in mixtures without any Caco-2 protein (volume replaced by assay buffer), control values (corresponding to 100 % activity) were obtained in mixtures with no substance added. The potency of the test substances in question, expressed as IC₅₀ values, was

calculated from dosage/activity curves consisting of 11 measuring points in each case.

The results obtained are shown in the following Table:

Compound (Example No.)	DPP IV inhibition IC ₅₀ (nM)
1	11
1(1)	24
1(2)	42
1(3)	110
1(4)	58
1(5)	134
1(6)	48
1(7)	434
1(8)	213
1(9)	61
1(11)	54
1(12)	18
1(13)	152
1(14)	158
1(15)	58
1(22)	48

The compounds prepared according to the invention are well tolerated, as for example when 30 mg/kg of the compound of Example 1 were administered to rats by oral route no toxic side effects or changes in the animals' behaviour could be detected.

In view of their ability to inhibit DPP-IV activity, the compounds of general formula I according to the invention and the corresponding pharmaceutically acceptable salts thereof are suitable for treating all those conditions or illnesses which can be influenced by the inhibition of the DPP-IV activity. It is

therefore to be expected that the compounds according to the invention will be suitable for the prevention or treatment of diseases or conditions such as type I and type II diabetes mellitus, diabetic complications, metabolic acidosis or ketosis, insulin resistance, dyslipidaemias of various origins, arthritis, atherosclerosis and related diseases, obesity, allograft transplantation and calcitonin-induced osteoporosis. In addition these substances are capable of preventing B-cell degeneration such as e.g. apoptosis or necrosis of pancreatic B-cells. The substances are also suitable for improving or restoring the function of pancreatic cells and also increasing the number and size of pancreatic B-cells. Additionally, and on the basis of the role of the Glucagon-Like Peptides, such as e.g. GLP-1 and GLP-2 and their link with DPP-IV inhibition, it is likely that the compounds according to the invention are suitable for achieving, inter alia, a sedative or anxiety-relieving effect and also favourably affecting catabolic states after operations or hormonal stress responses or of reducing mortality and morbidity after myocardial infarct. They are also suitable for treating all conditions which are connected with the abovementioned effects and which are mediated by GLP-1 or GLP-2. The compounds according to the invention may also be used as diuretics or antihypertensives and are suitable for preventing and treating acute renal failure. They are also suitable for the prevention and treatment of chronic inflammatory intestinal diseases. It is also expected that DPP-IV inhibitors and hence also the compounds according to the invention may be used to treat infertility or to improve fertility in humans or mammals, particularly when the infertility is connected with insulin resistance or polycystic ovary syndrome. The substances are also suitable for treating deficiencies of growth hormone which are associated with reduced stature.

The compounds according to the invention may also be used in conjunction with other active substances. Therapeutic agents which are suitable for such combinations include, for example, antidiabetics, such as metformin, sulphonylureas (e.g. glibenclamid, tolbutamide, glimepiride), nateglinide, repaglinide, thiazolidinedione (e.g. rosiglitazone, pioglitazone), PPAR-gamma agonists (e.g. GI 262570), alpha-glucosidase inhibitors (e.g. acarbose, voglibose), alpha2 antagonists, insulin and insulin analogues, GLP-1 and

GLP-1 analogues (e.g. exendin-4) or amylin. Also, inhibitors of protein tyrosine phosphatase 1, substances which influence deregulated glucose production in the liver, such as e.g. inhibitors of glucose-6-phosphatase, or fructose-1,6-bisphosphatase, glycogen phosphorylase, glucagon receptor antagonists and inhibitors of phosphoenol pyruvate carboxykinase, glycogen synthase kinase or pyruvate dehydrokinase, lipid lowering agents, such as HMG-CoA-reductase inhibitors (e.g. simvastatin, atorvastatin), fibrates (e.g. bezafibrate, fenofibrate), nicotinic acid and its derivatives, cholesterol resorption inhibitors such as for example ezetimibe, bile acid-binding substances such as for example cholestyramine, HDL-raising compounds such as for example inhibitors of CETP or regulators of ABC1 or active substances for the treatment of obesity, such as e.g. sibutramine or tetrahydrolipostatin or β_3 -agonists such as SB-418790 or AD-9677.

It is also possible to combine the compounds with drugs for treating high blood pressure such as e.g. α_1 antagonists or ACE inhibitors, diuretics, β -blockers, etc., or combinations thereof.

The dosage required to achieve such an effect is expediently, by intravenous route, 1 to 100 mg, preferably 1 to 30 mg, and by oral route 1 to 1000 mg, preferably 1 to 100 mg, in each case 1 to 4 times a day. For this purpose, the compounds of formula I prepared according to the invention, optionally combined with other active substances, may be incorporated together with one or more inert conventional carriers and/or diluents, e.g. with corn starch, lactose, glucose, microcrystalline cellulose, magnesium stearate, polyvinylpyrrolidone, citric acid, tartaric acid, water, water/ethanol, water/glycerol, water/sorbitol, water/polyethylene glycol, propylene glycol, cetylstearyl alcohol, carboxymethylcellulose or fatty substances such as hard fat or suitable mixtures thereof into conventional galenic preparations such as plain or coated tablets, capsules, powders, suspensions or suppositories.

The Examples that follow are intended to illustrate the invention:

Preparation of the starting compounds:

Example I

Ethyl 5-amino-3-benzyl-2-(3-tert.-butyloxycarbonylamino-piperidin-1-yl)-3H-imidazol-4-carboxylate

50 g (0.199 mol) of N-benzyl-N'-cyano-O-phenyl-isourea and 40.056 g (0.2 mol) of 3-tert.-butyloxycarbonylamino-piperidine are heated to 80°C in 50 ml of dimethylformamide (DMF) for 4 hours. After standing overnight 250 ml of ethyl acetate are added, the precipitate is suction filtered, washed with ethyl acetate and ether and dried. More product is obtained from the mother liquor after evaporation and treatment of the residue with ethyl acetate and ether. Total yield: 48.0 g (67.5% of theory) of N-benzyl-N'-cyano-(3-tert.-butyloxycarbonylamino-piperidin)-1-carboxamide.

R_f value: 0.56 (aluminium oxide, methylene chloride/methanol = 40:1)

10.008 g (28 mmol) of this substance are dissolved in 15 ml DMF. After the addition of 4.256 g (30.8 mmol) of potassium carbonate the mixture is treated with ultrasound for three minutes, then 3.416 ml (30.8 mmol) of ethyl bromoacetate are added in one go and the mixture is stirred for 36 hours at ambient temperature, adding another 10 ml of DMF after 8 hours for ease of stirring. The reaction mixture is stirred with water, extracted with ethyl acetate, the organic phase is dried and concentrated by evaporation. The resin obtained is purified by column chromatography (silica gel, ethyl acetate/petroleum ether = 3:1 to 9:1)

5.4 g (43.5 % of theory) of N-benzyl-N'-cyano-N-ethoxycarbonylmethyl-(3-tert.-butyloxy-carbonylamino-piperidin)-1-carboxamide are obtained.

R_f value: 0.7 (silica gel, ethyl acetate/petroleum ether = 4:1)

5.1 g (11.498 mmol) of this compound are added batchwise to a solution of 0.785 g (11.536 mmol) of sodium ethoxide in 25 ml of ethanol. The mixture is stirred for 40 minutes at 60°C, then combined with 50 ml of ethanol and 10 ml of water and cooled. The precipitate is suction filtered and dissolved in

methylene chloride. After drying and evaporation of the solvent, 4.8 g (94.1 % of theory) of the title compound is obtained.

R_f value: 0.4 (silica gel, ethyl acetate/petroleum ether = 4:1)

The following was obtained analogously to Example I:

(1) ethyl 5-amino-3-(3-methyl-but-2-enyl)-2-(3-tert.-butyloxycarbonylamino-piperidin-1-yl)-3H-imidazol-4-carboxylate

prepared from diphenyl-N-cyanocarbonimidate, glycinethylester, (3-tert.-butyloxy-carbonylamino)-piperidine and 3-methyl-but-2-en-1-yl-bromide.

R_f value: 0.1-0.2 (silica gel, ethyl acetate/petroleum ether = 1:1)

Example II

tert. butyl [1-(7-benzyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]carbamate

360 mg (0.812 mmol) of ethyl 5-amino-3-benzyl-2-(3-tert.-butyloxy-carbonylamino-piperidin-1-yl)-3H-imidazol-4-carboxylate and 131.472 (1.2 mmol) of ethylformidate hydrochloride are placed in 1.3 g phenol. A solution of 141.667 mg (2mmol) of sodiummethoxide in 2 ml of tetrahydrofuran (THF) is added dropwise with stirring, the THF is evaporated off and the mixture is kept for 2 hours at 150°C. The brown reaction mixture is purified through a silica gel column. 50 mg (15.7 % of theory) of the title compound were obtained. melting point: 208°C.

R_f value: 0.15 (silica gel, methylene chloride/methanol = 15:1)

Example III

tert. butyl [1-(7-benzyl-1-methyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]carbamate

19 mg (0.138 mmol) of potassium carbonate and then 16 mg (0.113 mmol) of methyl iodide are added to a solution of 42 mg (0.099 mmol) of the compound of Example II in 0.3 ml DMF. The mixture is stirred for 2 hours at ambient temperature, then triturated with water and extracted with ethyl acetate. The organic phase is washed with water and dried with activated

charcoal and magnesium sulphate. After evaporation 29 mg (66.8 % of theory) of the title compound is obtained.

R_f value: 0.3 (silica gel, methylene chloride/methanol = 15:1)

The following was obtained analogously to Example III:

(1) tert. butyl [1-(7-benzyl-1-{2-oxo-2-phenyl-ethyl}-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

prepared from Example II and phenacylbromide.

R_f value: 0.4 (silica gel, methylene chloride/methanol = 15:1)

Example IV

tert. butyl [1-(7-benzyl-2-methylsulphanyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

3.186 ml (27 mmol) of ethyloxycarbonyl isothiocyanate are added dropwise to a solution of 10.8 g (24.345 mmol) of the compound of Example I in 45 ml THF. The mixture is heated to boiling for one hour, concentrated by evaporation and the residue is brought to crystallisation by treating with diisopropylether.

13.5 g (96.5 % of theory) of ethyl 3-benzyl-2-(3-tert.-butyloxycarbonylamino-piperidin-1-yl)-5-(N'-ethyloxycarbonyl-thioureido)-3H-imidazol-4-carboxylate are obtained.

13 g (22.620 mmol) of this compound are placed in 21 ml of n-butanol. After the addition of 2.536 (22.6 mmol) of potassium-tert.-butoxide the mixture is stirred for 45 minutes at 100°C, during which time a precipitate settles out. After standing overnight at ambient temperature the mixture is combined with ether, suction filtered and dried. 10.6 g (94.7 % of theory) of tert. butyl [1-(7-benzyl-2-mercapto-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate potassium salt are obtained.

10.5 g (21.225 mmol) of this compound are suspended in 25 ml of water, and ethanol is added to the solution. After the addition of 2.135 ml (21.515 mmol) of dimethylsulphate the mixture is stirred for 4 hours at ambient temperature.

The precipitate is suction filtered, washed with cold ethanol and dried. 8.8 g (88.1 % of theory) of the title compound are obtained.

R_f value: 0.3 (silica gel, methylene chloride/methanol = 20:1)

The following compounds were obtained analogously to Example IV:

(1) tert. butyl [1-(7-benzyl-2-benzylsulphanyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

prepared from compound I, ethyloxycarbonyl isothiocyanate and benzylbromide.

R_f value: 0.65 (silica gel, ethyl acetate/petroleum ether = 2:1)

(2) tert. butyl [1-(7-allyl-2-methylsulphanyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

prepared from compound I.1, ethyloxycarbonyl isothiocyanate and methyl iodide.

R_f value: 0.6 (silica gel, methylene chloride/methanol = 10:1)

(3) tert. butyl [1-(7-benzyl-2-[2-phenylethyl]sulphanyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

prepared from compound I, ethyloxycarbonyl isothiocyanate and 2-phenylethylbromide.

R_f value: 0.65 (silica gel, ethyl acetate/petroleum ether = 2:1)

Example V

tert-butyl [1-(7-benzyl-1-methyl-2-methylsulphanyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

A suspension of 2.5 g (5.312 mmol) of compound IV in 15 ml DMF is combined with 645.15 mg (5.75 mmol) of potassium-tert.-butoxide. 922.605 mg (6.5 mmol) of methyl iodide are added to the resulting solution and the mixture is stirred overnight at ambient temperature. Water is added and the mixture is extracted with methylene chloride. The organic phase is washed with water, dried and concentrated by evaporation. The residue is crystallised

with diisopropylether. 2 g (77.7 % of theory) of the title compound are obtained.

R_f value: 0.55 (silica gel, methylene chloride/methanol = 20:1)

The following compounds were obtained analogously to Example V:

(1) tert. butyl [1-(7-benzyl-2-methylsulphanyl-1-phenacyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminat

prepared from compound IV and phenacylbromide.

R_f value: 0.7 (aluminium oxide, methylene chloride/methanol = 40:1)

(2) tert. butyl [1-(1-methyl-7-(3-methyl-butenyl)-2-methylsulphanyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminat

prepared from compound IV.1 and methyl iodide.

R_f value: 0.4 (silica gel, methylene chloride/methanol = 20:1)

(3) tert. butyl [1-(1-benzyl-7-benzyl-2-methylsulphanyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminat

prepared from compound IV and benzylbromide.

R_f value: 0.75 (silica gel, ethyl acetate/petroleum ether = 4:1)

(4) tert. butyl [1-(7-benzyl-2-methylsulphanyl-1-(2-phenylethyl)-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminat

prepared from compound IV and 2-phenylethylbromide.

R_f value: 0.75 (silica gel, ethyl acetate/petroleum ether = 4:1)

(5) tert. butyl [1-(7-benzyl-2-benzylsulphanyl-1-methyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminat

prepared from compound IV.1 and methyl iodide.

R_f value: 0.85 (silica gel, ethyl acetate/petroleum ether = 1:2)

(6) tert. butyl [1-(7-benzyl-2-[2-phenylethyl]sulphanyl-1-methyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminat

prepared from compound IV.3 and methyl iodide.

R_f value: 0.65 (silica gel, ethyl acetate/petroleum ether = 1:2)

Example VI

tert. butyl [1-(7-benzyl-1-methyl-2-methylsulphinyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate and tert. butyl [1-(7-benzyl-1-methyl-2-methylsulphonyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

A solution of 250 mg (0.516 mmol) of compound V in 5 ml of dichloromethane and 0.5 ml of methanol is combined with 120.789 mg (0.7 mmol) of m-chloro-perbenzoic acid with stirring and cooling with ice. After 30 minutes the ice bath is removed and the mixture is stirred overnight at ambient temperature. 50 ml of methylene chloride are added and the mixture is extracted with 10 % soda solution. The organic phase is washed with water, dried and concentrated by evaporation. 220 mg of the two title compounds are obtained in the ratio 45 : 55.

R_f value: 0.1 (sulphoxide) and 0.8 (sulphone) (silica gel, ethyl acetate)

The following was obtained analogously to Example VI:

(1) tert. butyl [1-methyl-7-(3-methyl-but-2-enyl)-2-methylsulphinyl-6-oxo-6,7-dihydro-1H-purin-8-yl]-piperidin-3-yl)-carbaminate and tert.butyl [1-methyl-7-(3-methyl-but-2-enyl)-2-methylsulphonyl-6-oxo-6,7-dihydro-1H-purin-8-yl-]-piperidin-3-yl)-carbaminate

prepared from compound V.2 and m-chloro-perbenzoic acid. The product obtained after a reaction time of 80 minutes is the sulphoxide, which contains a maximum of 10% sulphone.

R_f value: 0.5 (silica gel, methylene chloride/methanol = 10:1)

Example VII

Tert. butyl [1-(7-benzyl –2-benzylamino-1-methyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

258.312 mg of the mixture obtained in Example VI and 214.313 mg of benzylamine are stirred for 16 hours at ambient temperature. The mixture is

trituated with 20 ml of diisopropylether, the precipitate is suction filtered, dissolved in a little methylene chloride and crystallised with diisopropylether. 250 mg of the title compound are obtained.

R_f value: 0.55 (silica gel, ethyl acetate)

The following compounds were prepared analogously to Example VII:

(1) tert. butyl [1-(7-benzyl-2-[4-fluoro-benzyl]amino-1-methyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

prepared from Example VI and 4-fluoro-benzylamine

R_f value: 0.49 (silica gel, ethyl acetate)

(2) tert. butyl [1-(7-benzyl-1-methyl-2-(2-phenylethyl)amino-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

prepared from Example VI and 2-(2-phenylethyl)amine.

R_f value: 0.5 (silica gel, ethyl acetate)

(3) tert. butyl [1-(7-benzyl-2-isopropylamino-1-methyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

prepared from Example VI and isopropylamine.

R_f value: 0.6 (silica gel, methylene chloride/methanol = 9:1)

(4) tert. butyl [1-(7-benzyl-1-methyl-2-methylamino-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

prepared from Example VI and methylamine gas.

R_f value: 0.27 (silica gel, methylene chloride/methanol = 19:1)

(5) tert. butyl [1-(7-benzyl-2-cyclohexylamino-1-methyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

prepared from Example VI and cyclohexylamine

R_f value: 0.65 (silica gel, methylene chloride/methanol = 9:1)

(6) tert. butyl [1-(7-benzyl-2-cyclohexylmethylamino-1-methyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

prepared from Example VI and cyclohexylmethanamine.

R_f value: 0.54 (silica gel, ethyl acetate)

(7) tert. butyl [1-(7-benzyl-1-methyl-2-piperidino-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

prepared from Example VI and piperidine.

R_f value: 0.45 (silica gel, methylene chloride/methanol = 20:1)

(8) tert. butyl [1-(7-benzyl-2-dimethylamino-1-methyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

prepared from Example VI and dimethylamine

R_f value: 0.65 (silica gel, methylene chloride/methanol = 9:1)

(9) tert. butyl [1-(2-amino-7-benzyl-1-methyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

prepared from Example VI and ammonia gas.

R_f value: 0.4 (silica gel, methylene chloride/methanol = 10:1)

(10) tert. butyl [1-(2-benzylamino-1-methyl-7-[3-methylbut-2-enyl]-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidino-3-yl]-carbaminate

prepared from Example VI.1 and benzylamine

R_f value: 0.6 (silica gel, ethyl acetate)

Example VIIITert. butyl [1-(7-benzyl-1-methyl-2-methyl-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

A solution of 258.32 mg of Example VI in 3 ml of THF is combined with 0.2 ml of a 3-molar solution of methylmagnesium bromide in ether and the mixture is stirred for 48 hours at ambient temperature. 50 ml of ether were added and the mixture was extracted with water at pH 4. The organic phase was dried and concentrated by evaporation. The product obtained was purified through a silica gel column. 55 mg of the title compound are obtained.

R_f value: 0.55 (silica gel, methylene chloride/methanol = 10:1)

The following compound was obtained analogously to Example VIII:

(1) tert. butyl [1-(1-methyl-7-[3-methyl-but-2-enyl]-2-[2-phenylethyl]-6-oxo-6,7-dihydro-1H-purin-8-yl)-piperidin-3-yl]-carbaminate

prepared from Example VI.1 and (2-phenylethyl)magnesium bromide.

R_f value: 0.5 (silica gel; methylene chloride/methanol = 10:1)

Examples of the preparation of the end products:

Example 18-(3-amino-piperidin-1-yl)-7-benzyl-2-benzylamino-1-methyl-1,7-dihydro-purin-6-one

A solution of 200 mg of Example VII in 2 ml of dichloromethane is combined with 3 ml of trifluoroacetic acid with stirring and cooling with ice. After 30 minutes the ice bath is removed and stirring is continued for 2 hours. The solvent is evaporated at low temperature, the residue is triturated with ether, suction filtered and dried in vacuo. 150 mg (73.1 % of theory) of the trifluoroacetate of the title compound are obtained.

R_f value: 0.45 (aluminium oxide, methylene chloride/methanol = 20:1)

¹H-NMR spectrum (400 MHz, DMSO-d₆):

1.5 (m,2H), 1,7 (m,1H), 1.95 (m,1H), 2.8 (m,1H), 3.0 (m,1H), 3.15 (d,1H), 3.4 (s+m,4H), 3.5(d,1H), 4.55 (d,2H), 5.35(s,2H), 7.1-7.5 (m,12H), 8.0 (s,3H), 8.1-8.3 (m,1H)

The following compounds were obtained analogously to Example 1:

(1) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-(4-fluoro-benzylamino)-1-methyl-1,7-dihydro-purin-6-one trifluoroacetate

prepared from Example VII.1 and trifluoroacetic acid

R_f value: 0.69 (aluminium oxide, methylene chloride/methanol = 9:1)

(2) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(2-phenylethyl)amino]-1,7-dihydro-purin-6-one trifluoroacetate

prepared from Example VII.2 and trifluoroacetic acid.

R_f value: 0.75 (aluminium oxide, methylene chloride/methanol = 9:1)

(3) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-isopropylamino-1-methyl-1,7-dihydro-purin-6-one trifluoroacetate

prepared from Example VII.3 and trifluoroacetic acid.

R_f value: 0.68 (aluminium oxide, methylene chloride/methanol = 9:1)

(4) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-methylamino-1,7-dihydro-purin-6-one trifluoroacetate

prepared from Example VII.4 and trifluoroacetic acid.

R_f value: 0.26 (aluminium oxide, methylene chloride/methanol = 9:1)

(5) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-cyclohexylamino-1-methyl-1,7-dihydro-purin-6-one trifluoroacetate

prepared from Example VII.5 and trifluoroacetic acid.

R_f value: 0.65 (aluminium oxide, methylene chloride/methanol = 9:1)

(6) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-cyclohexylmethylamino-1-methyl-1,7-dihydro-purin-6-one trifluoroacetate

prepared from Example VII.6 and trifluoroacetic acid.

R_f value: 0.69 (aluminium oxide, methylene chloride/methanol = 9:1)

(7) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(piperidin-1-yl)-1,7-dihydro-purin-6-one hydrochloride

prepared from Example VII.7 and hydrogen chloride in dioxane.

R_f value: 0.3-0.5 (aluminium oxide, methylene chloride/methanol = 20:1)

(8) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-dimethylamino-1-methyl-1,7-dihydro-purin-6-one trifluoroacetate

prepared from Example VII.8 and trifluoroacetic acid.

R_f value: 0.69 (aluminium oxide, methylene chloride/methanol = 9:1)

(9) 2-amino-8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-1,7-dihydro-purin-6-one trifluoroacetate

prepared from Example VII.9 and trifluoroacetic acid.

R_f value: 0.2 (aluminium oxide, methylene chloride/methanol = 10:1)

(10) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(3-methyl-but-2-en-1-yl)-1,7-dihydro-purin-6-one trifluoroacetate

prepared from Example VII.10 and trifluoroacetic acid.

R_f value: 0.55 (aluminium oxide, methylene chloride/methanol = 10:1)

(11) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-methyl-1,7-dihydro-purin-6-one trifluoroacetate

prepared from Example VIII and trifluoroacetic acid.

R_f value: 0.6 (aluminium oxide, methylene chloride/methanol = 10:1)

(12) 8-(3-amino-piperidin-1-yl)-1-methyl-7-(3-methyl-but-2-en-1-yl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one trifluoroacetate

prepared from Example VIII.1 and trifluoroacetic acid.

R_f value: 0.4 (aluminium oxide, methylene chloride/methanol = 20:1)

(13) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-1,7-dihydro-purin-6-one hydrochloride

prepared from Example III and hydrogen chloride in dioxane.

R_f value: 0.2-0.5 (aluminium oxide, methylene chloride/methanol = 20:1)

(14) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-methylsulphanyl-1,7-dihydro-purin-6-one hydrochloride

prepared from Example IV and hydrogen chloride in dioxane.

R_f value: 0.5 (aluminium oxide, methylene chloride/methanol = 10:1)

(15) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-benzylsulphanyl-1,7-dihydro-purin-6-one trifluoroacetate

prepared from Example IV.1 and trifluoroacetic acid.

R_f value: 0.5-0.6 (aluminium oxide, methylene chloride/methanol = 10:1)

(16) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-methylsulphanyl-1,7-dihydro-purin-6-one hydrochloride

prepared from Example V and hydrogen chloride in dioxane.

R_f value: 0.5 (aluminium oxide, methylene chloride/methanol = 20:1)

(17) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-methylsulphanyl-1-(2-phenyl-2-oxoethyl)-1,7-dihydro-purin-6-one hydrochloride

prepared from Example V.1 and hydrogen chloride in dioxane.

R_f value: 0.4 (aluminium oxide, methylene chloride/methanol = 20:1)

(18) 8-(3-amino-piperidin-1-yl)-1-benzyl-7-benzyl-2-methylsulphanyl-1,7-dihydro-purin-6-one hydrochloride

prepared from Example V.3 and hydrogen chloride.

R_f value: 0.4-0.5 (aluminium oxide, methylene chloride/methanol = 20:1)

(19) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-methylsulphanyl-1-(2-phenylethyl)-1,7-dihydro-purin-6-one hydrochloride

prepared from Example V.4 and hydrogen chloride.

R_f value: 0.4 (aluminium oxide, methylene chloride/methanol = 20:1)

(20) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-benzylsulphanyl-1-methyl-1,7-dihydro-purin-6-one

prepared from Example V.5 and hydrogen chloride.

R_f value: 0.5 (aluminium oxide, methylene chloride/methanol = 20:1)

(21) 8-(3-amino-piperidin-1-yl)-7-benzyl-2-[(2-phenylethyl)sulphanyl]-1-methyl-1,7-dihydro-purin-6-one

prepared from Example V.6 and hydrogen chloride.

R_f value: 0.55 (aluminium oxide, methylene chloride/methanol = 20:1)

(22) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2-oxo-2-phenyl-ethyl)-1,7-dihydro-purin-6-one trifluoroacetate

prepared from Example III.1 and trifluoroacetic acid.

R_f value: 0.65 (aluminium oxide, methylene chloride/methanol = 10:1)

The following compounds may also be obtained analogously to the preceding Examples and other methods known from the literature:

(1) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-ethyl-2-benzylamino-1,7-dihydro-purin-6-one

(2) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-propyl-2-benzylamino-1,7-dihydro-purin-6-one

(3) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-isopropyl-2-benzylamino-1,7-dihydro-purin-6-one

(4) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-butyl-2-benzylamino-1,7-dihydro-purin-6-one

(5) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2-methyl-propyl)-2-benzylamino-1,7-dihydro-purin-6-one

- (6) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-pentyl-2-benzylamino-1,7-dihydro-purin-6-one
- (7) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(3-methyl-butyl)-2-benzylamino-1,7-dihydro-purin-6-one
- (8) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-hexyl-2-benzylamino-1,7-dihydro-purin-6-one
- (9) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(4-methyl-pentyl)-2-benzylamino-1,7-dihydro-purin-6-one
- (10) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-allyl-2-benzylamino-1,7-dihydro-purin-6-one
- (11) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(3-methyl-but-2-en-1-yl)-2-benzylamino-1,7-dihydro-purin-6-one
- (12) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(prop-2-in-1-yl)-2-benzylamino-1,7-dihydro-purin-6-one
- (13) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-cyclopropylmethyl-2-benzylamino-1,7-dihydro-purin-6-one
- (14) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-cyclohexylmethyl-2-benzylamino-1,7-dihydro-purin-6-one
- (15) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(4-chlorobenzyl)-2-benzylamino-1,7-dihydro-purin-6-one
- (16) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2-fluorobenzyl)-2-benzylamino-1,7-dihydro-purin-6-one

(17) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(4-methylbenzyl)-2-benzylamino-1,7-dihydro-purin-6-one

(18) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2,4-dimethylbenzyl)-2-benzylamino-1,7-dihydro-purin-6-one

(19) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2,4-dimethoxybenzyl)-2-benzylamino-1,7-dihydro-purin-6-one

(20) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(3-trifluoromethyl-benzyl)-2-benzylamino-1,7-dihydro-purin-6-one

(21) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(3-trifluoromethoxy-benzyl)-2-benzylamino-1,7-dihydro-purin-6-one

(22) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(3-difluoromethoxy-benzyl)-2-benzylamino-1,7-dihydro-purin-6-one

(23) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(furan-2-yl-methyl)-2-benzylamino-1,7-dihydro-purin-6-one

(24) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(thien-2-yl-methyl)-2-benzylamino-1,7-dihydro-purin-6-one

(25) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(3-methyl-isoxazol-5-yl-methyl)-2-benzylamino-1,7-dihydro-purin-6-one

(26) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(pyridin-4-yl-methyl)-2-benzylamino-1,7-dihydro-purin-6-one

(27) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(4-methyl-phenyl)-2-oxo-ethyl]-2-benzylamino-1,7-dihydro-purin-6-one

- (28) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(2,4-dichloro-phenyl)-2-oxo-ethyl]-2-benzylamino-1,7-dihydro-purin-6-one
- (29) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(4-methoxy-phenyl)-2-oxo-ethyl]-2-benzylamino-1,7-dihydro-purin-6-one
- (30) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(4-trifluoromethyl-phenyl)-2-oxo-ethyl]-2-benzylamino-1,7-dihydro-purin-6-one
- (31) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(4-trifluoromethoxy-phenyl)-2-oxo-ethyl]-2-benzylamino-1,7-dihydro-purin-6-one
- (32) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(4-difluoromethoxy-phenyl)-2-oxo-ethyl]-2-benzylamino-1,7-dihydro-purin-6-one
- (33) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(furan-2-yl)-2-oxo-ethyl]-2-benzylamino-1,7-dihydro-purin-6-one
- (34) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(thien-2-yl)-2-oxo-ethyl]-2-benzylamino-1,7-dihydro-purin-6-one
- (35) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(pyridin-3-yl)-2-oxo-ethyl]-2-benzylamino-1,7-dihydro-purin-6-one
- (36) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2-oxopropyl)-2-benzylamino-1,7-dihydro-purin-6-one
- (37) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2-cyclohexyl-2-oxo-ethyl)-2-benzylamino-1,7-dihydro-purin-6-one
- (38) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-ethyl-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(39) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-propyl-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(40) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-isopropyl-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(41) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-butyl-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(42) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2-methyl-propyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(43) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-pentyl-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(44) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(3-methyl-butyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(45) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-hexyl-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(46) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(4-methyl-pentyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(47) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-allyl-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(48) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(3-methyl-but-2-en-1-yl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(49) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(prop-2-in-1-yl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

- (50) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-cyclopropylmethyl-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (51) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-cyclohexylmethyl-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (52) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(4-chlorobenzyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (53) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2-fluorobenzyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (54) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(4-methylbenzyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (55) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2,4-dimethylbenzyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (56) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2,4-dimethoxybenzyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (57) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(3-trifluoromethyl-benzyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (58) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(3-trifluoromethoxy-benzyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (59) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(3-difluoromethoxy-benzyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (60) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(furan-2-yl-methyl)-2-(2-phenylethyl)-7-dihydro-purin-6-one

- (61) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(thien-2-yl-methyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (62) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(3-methyl-isoxazol-5-yl-methyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (63) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(pyridin-4-yl-methyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (64) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(4-methyl-phenyl)-2-oxo-ethyl]-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (65) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(2,4-dichloro-phenyl)-2-oxo-ethyl]-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (66) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(4-methoxy-phenyl)-2-oxo-ethyl]-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (67) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(4-trifluoromethyl-phenyl)-2-oxo-ethyl]-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (68) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(4-trifluoromethoxy-phenyl)-2-oxo-ethyl]-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (69) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(4-difluoromethoxy-phenyl)-2-oxo-ethyl]-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (70) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(furan-2-yl)-2-oxo-ethyl]-2-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (71) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(thien-2-yl)-2-oxo-ethyl]-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(72) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(pyridin-3-yl)-2-oxo-ethyl]-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(73) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2-oxo-propyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(74) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2-cyclohexyl-2-oxo-ethyl)-2-(2-phenylethyl)-1,7-dihydro-purin-6-one

(75) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-ethylamino-1,7-dihydro-purin-6-one

(76) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-propylamino-1,7-dihydro-purin-6-one

(77) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-butylamino-1,7-dihydro-purin-6-one

(78) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(2-methyl-propylamino)-1,7-dihydro-purin-6-one

(79) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(N-ethyl-N-methyl-amino)-1,7-dihydro-purin-6-one

(80) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-diethylamino-1,7-dihydro-purin-6-one

(81) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(N-methyl-N-propyl-amino)-1,7-dihydro-purin-6-one

(82) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(N-butyl-N-methyl-amino)-1,7-dihydro-purin-6-one

- (83) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(2-hydroxyethyl-amino)-1,7-dihydro-purin-6-one
- (84) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[N-methyl-N-(2-hydroxyethyl)-amino]-1,7-dihydro-purin-6-one
- (85) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(2-methoxy-ethylamino)-1,7-dihydro-purin-6-one
- (86) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(3-methoxy-propylamino)-1,7-dihydro-purin-6-one
- (87) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[N-methyl-N-(2-methoxy-ethyl)-amino]-1,7-dihydro-purin-6-one
- (88) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(2-phenoxyethyl)-amino]-1,7-dihydro-purin-6-one
- (89) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(2-aminoethyl)-amino]-1,7-dihydro-purin-6-one
- (90) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(3-aminopropyl)-amino]-1,7-dihydro-purin-6-one
- (91) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(2-methylamino-ethyl)-amino]-1,7-dihydro-purin-6-one
- (92) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(3-dimethylamino-propyl)amino]-1,7-dihydro-purin-6-one
- (93) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(2-acetyl-amino-ethyl)amino]-1,7-dihydro-purin-6-one

- (94) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(2-benzoylamino-ethyl)amino]-1,7-dihydro-purin-6-one
- (95) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(2-ethoxycarbonylamino-ethyl)amino]-1,7-dihydro-purin-6-one
- (96) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(2-methylsulphanyl-ethyl)amino]-1,7-dihydro-purin-6-one
- (97) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(2-phenylsulphanyl-ethyl)amino]-1,7-dihydro-purin-6-one
- (98) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(2-cyano-ethyl)amino]-1,7-dihydro-purin-6-one
- (99) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(carboxymethyl)amino]-1,7-dihydro-purin-6-one
- (100) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(ethoxycarbonylmethyl)amino]-1,7-dihydro-purin-6-one
- (101) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(aminocarbonylmethyl)amino]-1,7-dihydro-purin-6-one
- (102) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(ethylaminocarbonylmethyl)amino]-1,7-dihydro-purin-6-one
- (103) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(dimethylaminocarbonylmethyl)amino]-1,7-dihydro-purin-6-one
- (104) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(pyrrolidin-1-yl-carbonylmethyl)amino]-1,7-dihydro-purin-6-one

(105) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(piperidin-1-yl-carbonylmethyl)amino]-1,7-dihydro-purin-6-one

(106) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(morpholin-4-ylcarbonylmethyl)amino]-1,7-dihydro-purin-6-one

(107) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(cyclopentylmethyl)amino]-1,7-dihydro-purin-6-one

(108) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(cyclobutylmethyl)amino]-1,7-dihydro-purin-6-one

(109) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(cyclopropylmethyl)amino]-1,7-dihydro-purin-6-one

(110) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(2-cyclohexylethyl)amino]-1,7-dihydro-purin-6-one

(111) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(3-cyclohexylpropyl)amino]-1,7-dihydro-purin-6-one

(112) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(cyclobutylamino)-1,7-dihydro-purin-6-one

(113) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(cyclopentylamino)-1,7-dihydro-purin-6-one

(114) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(phenylamino)-1,7-dihydro-purin-6-one

(115) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(3,4-dichlorobenzylamino)-1,7-dihydro-purin-6-one

(116) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(4-methylbenzyl-amino)-1,7-dihydro-purin-6-one

(117) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(4-methoxybenzyl-amino)-1,7-dihydro-purin-6-one

(118) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(4-trifluoromethylbenzyl-amino)-1,7-dihydro-purin-6-one

(119) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(4-trifluoromethoxybenzyl-amino)-1,7-dihydro-purin-6-one

(120) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(4-difluoromethoxybenzyl-amino)-1,7-dihydro-purin-6-one

(121) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(3,4-methylenedioxybenzyl-amino)-1,7-dihydro-purin-6-one

(122) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(furan-2-yl-methyl)amino]-1,7-dihydro-purin-6-one

(123) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(thien-2-yl-methyl)amino]-1,7-dihydro-purin-6-one

(124) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(pyridine-2-yl-methyl)amino]-1,7-dihydro-purin-6-one

(125) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(4-methylthiazol-2-ylmethyl)amino]-1,7-dihydro-purin-6-one

(126) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[[2-(pyridine-2-yl)ethyl]amino]-1,7-dihydro-purin-6-one

- (127) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(pyrrolidin-1-yl)-1,7-dihydro-purin-6-one
- (128) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(morpholin-4-yl)-1,7-dihydro-purin-6-one
- (129) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(piperazin-1-yl)-1,7-dihydro-purin-6-one
- (130) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(4-methyl-piperazin-1-yl)-1,7-dihydro-purin-6-one
- (131) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-ethyl-1,7-dihydro-purin-6-one
- (132) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-propyl-1,7-dihydro-purin-6-one
- (133) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-isopropyl-1,7-dihydro-purin-6-one
- (134) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-butyl-1,7-dihydro-purin-6-one
- (135) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(2-methyl-propyl)-1,7-dihydro-purin-6-one
- (136) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-pentyl-1,7-dihydro-purin-6-one
- (137) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(3-methyl-butyl)-1,7-dihydro-purin-6-one

(138) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-hexyl-1,7-dihydro-purin-6-one

(139) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(4-methyl-pentyl)-1,7-dihydro-purin-6-one

(140) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-allyl-1,7-dihydro-purin-6-one

(141) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(3-methyl-but-2-en-1-yl)-1,7-dihydro-purin-6-one

(142) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(hex-5-en-1-yl)-1,7-dihydro-purin-6-one

(143) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(pent-4-en-1-yl)-1,7-dihydro-purin-6-one

(144) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(prop-2-yn-1-yl)-1,7-dihydro-purin-6-one

(145) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(2-phenylethenyl)-1,7-dihydro-purin-6-one

(146) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(4-methyl-phenyl)ethenyl]-1,7-dihydro-purin-6-one

(147) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(2-chloro-phenyl)ethenyl]-1,7-dihydro-purin-6-one

(148) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(3-fluoro-phenyl)ethenyl]-1,7-dihydro-purin-6-one

(149) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(3,4-dimethoxy-phenyl)ethenyl]-1,7-dihydro-purin-6-one

(150) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(3-chloro-phenyl)ethenyl]-1,7-dihydro-purin-6-one

(151) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(3-trifluoromethyl-phenyl)ethenyl]-1,7-dihydro-purin-6-one

(152) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(3-trifluoromethoxy-phenyl)ethenyl]-1,7-dihydro-purin-6-one

(153) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(3-phenyl-prop-2-en-1-yl)-1,7-dihydro-purin-6-one

(154) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-benzyl-1,7-dihydro-purin-6-one

(155) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-phenyl-1,7-dihydro-purin-6-one

(156) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(3-phenyl-propyl)-1,7-dihydro-purin-6-one

(157) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(4-fluoro-benzyl)-1,7-dihydro-purin-6-one

(158) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(3-trifluoromethyl-benzyl)-1,7-dihydro-purin-6-one

(159) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(3-chloro-benzyl)-1,7-dihydro-purin-6-one

(160) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(3-bromo-benzyl)-1,7-dihydro-purin-6-one

(161) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(3-trifluoromethoxy-benzyl)-1,7-dihydro-purin-6-one

(162) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(3-methoxy-benzyl)-1,7-dihydro-purin-6-one

(163) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(3-chloro-phenyl)ethyl]-1,7-dihydro-purin-6-one

(164) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(2-fluoro-phenyl)ethyl]-1,7-dihydro-purin-6-one

(165) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(4-methyl-phenyl)ethyl]-1,7-dihydro-purin-6-one

(166) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(4-fluoro-phenyl)ethyl]-1,7-dihydro-purin-6-one

(167) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(3-trifluoromethyl-phenyl)ethyl]-1,7-dihydro-purin-6-one

(168) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-cyclopentyl-1,7-dihydro-purin-6-one

(169) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-cyclohexyl-1,7-dihydro-purin-6-one

(170) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(cyclohexylmethyl)-1,7-dihydro-purin-6-one

(171) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(cyclopentylmethyl)-1,7-dihydro-purin-6-one

(172) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(2-cyclohexyl-ethyl)-1,7-dihydro-purin-6-one

(173) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(furan-2-yl)-1,7-dihydro-purin-6-one

(174) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(thien-2-yl)-1,7-dihydro-purin-6-one

(175) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(pyridin-2-yl)-1,7-dihydro-purin-6-one

(176) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(pyridin-4-yl)-1,7-dihydro-purin-6-one

(177) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(furan-2-yl-methyl)-1,7-dihydro-purin-6-one

(178) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(thien-2-yl-methyl)-1,7-dihydro-purin-6-one

(179) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-(pyridin-2-yl-methyl)-1,7-dihydro-purin-6-one

(180) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(furan-2-yl)ethyl]-1,7-dihydro-purin-6-one

(181) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(pyridin-2-yl)ethyl]-1,7-dihydro-purin-6-one

(182) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[3-(furan-2-yl)propyl]-1,7-dihydro-purin-6-one

(183) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[3-(pyridin-2-yl)propyl]-1,7-dihydro-purin-6-one

(184) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(3-chloro-but-2-en-1-yl)-1,7-dihydro-purin-6-one

(185) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(3-chloro-prop-2-en-1-yl)-1,7-dihydro-purin-6-one

(186) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(3-trifluoromethyl-3-chloro-prop-2-en-1-yl)-1,7-dihydro-purin-6-one

(187) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(3,3-dichloro-prop-2-en-1-yl)-1,7-dihydro-purin-6-one

(188) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(but-2-yn-1-yl)-1,7-dihydro-purin-6-one

(189) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(cyclohex-1-en-1-yl-methyl)-1,7-dihydro-purin-6-one

(190) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(cyclohexylmethyl)-1,7-dihydro-purin-6-one

(191) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(4-fluoro-benzyl)-1,7-dihydro-purin-6-one

(192) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(2-chloro-benzyl)-1,7-dihydro-purin-6-one

(193) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(3-bromo-benzyl)-1,7-dihydro-purin-6-one

(194) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(3-methyl-benzyl)-1,7-dihydro-purin-6-one

(195) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(3-trifluoromethyl-benzyl)-1,7-dihydro-purin-6-one

(196) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(2-cyano-benzyl)-1,7-dihydro-purin-6-one

(197) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(2-methoxy-benzyl)-1,7-dihydro-purin-6-one

(198) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(3-trifluoromethoxy-benzyl)-1,7-dihydro-purin-6-one

(199) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(3-difluoromethoxy-benzyl)-1,7-dihydro-purin-6-one

(200) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(3,4-difluoro-benzyl)-1,7-dihydro-purin-6-one

(201) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(naphth-1-yl-methyl)-1,7-dihydro-purin-6-one

(202) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(furan-2-yl-methyl)-1,7-dihydro-purin-6-one

(203) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(thien-2-yl-methyl)-1,7-dihydro-purin-6-one

- (204) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(pyridin-2-yl-methyl)-1,7-dihydro-purin-6-one
- (205) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(3-chloro-but-2-en-1-yl)-1,7-dihydro-purin-6-one
- (206) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(3-chloro-prop-2-en-1-yl)-1,7-dihydro-purin-6-one
- (207) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(3-trifluoromethyl-3-chloro-prop-2-en-1-yl)-1,7-dihydro-purin-6-one
- (208) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(3,3-dichloro-prop-2-en-1-yl)-1,7-dihydro-purin-6-one
- (209) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(but-2-yn-1-yl)-1,7-dihydro-purin-6-one
- (210) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(cyclohex-1-en-1-yl-methyl)-1,7-dihydro-purin-6-one
- (211) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(cyclohexylmethyl)-1,7-dihydro-purin-6-one
- (212) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(4-fluorobenzyl)-1,7-dihydro-purin-6-one
- (213) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(2-chlorobenzyl)-1,7-dihydro-purin-6-one
- (214) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(3-bromobenzyl)-1,7-dihydro-purin-6-one

- (215) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(3-methyl-benzyl)-1,7-dihydro-purin-6-one
- (216) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(3-trifluoromethyl-benzyl)-1,7-dihydro-purin-6-one
- (217) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(2-cyano-benzyl)-1,7-dihydro-purin-6-one
- (218) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(2-methoxy-benzyl)-1,7-dihydro-purin-6-one
- (219) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(3-trifluoromethoxy-benzyl)-1,7-dihydro-purin-6-one
- (220) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(3-difluoromethoxy-benzyl)-1,7-dihydro-purin-6-one
- (221) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(3,4-difluoro-benzyl)-1,7-dihydro-purin-6-one
- (222) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(naphth-1-yl-methyl)-1,7-dihydro-purin-6-one
- (223) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(furan-2-yl-methyl)-1,7-dihydro-purin-6-one
- (224) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(thien-2-yl-methyl)-1,7-dihydro-purin-6-one
- (225) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-(pyridin-2-yl-methyl)-1,7-dihydro-purin-6-one

- (226) 8-(3-amino-piperidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-benzyl-1,7-dihydro-purin-6-one
- (227) 8-(3-amino-pyrrolidin-1-yl)-2-benzylamino-1-methyl-7-benzyl-1,7-dihydro-purin-6-one
- (228) 8-(piperidin-3-yl)-2-benzylamino-1-methyl-7-benzyl-1,7-dihydro-purin-6-one
- (229) 8-(piperidin-4-yl)-2-benzylamino-1-methyl-7-benzyl-1,7-dihydro-purin-6-one
- (230) 8-(3-amino-hexahydroazepin-1-yl)-2-benzylamino-1-methyl-7-benzyl-1,7-dihydro-purin-6-one
- (231) 8-(4-amino-hexahydroazepin-1-yl)-2-benzylamino-1-methyl-7-benzyl-1,7-dihydro-purin-6-one
- (232) 8-(piperazin-1-yl)-2-benzylamino-1-methyl-7-benzyl-1,7-dihydro-purin-6-one
- (233) 8-(1,4-diazepan-1-yl)-2-benzylamino-1-methyl-7-benzyl-1,7-dihydro-purin-6-one
- (234) 8-(2-amino-cyclohexylamino)-2-benzylamino-1-methyl-7-benzyl-1,7-dihydro-purin-6-one
- (235) 8-(3-amino-cyclohexyl)-2-benzylamino-1-methyl-7-benzyl-1,7-dihydro-purin-6-one
- (236) 8-(3-amino-pyrrolidin-1-yl)-2-(2-phenylethyl)-1-methyl-7-benzyl-1,7-dihydro-purin-6-one

(237) 8-(piperidin-3-yl)-2-(2-phenylethyl)-1-methyl-7-benzyl-1,7-dihydro-purin-6-one

(238) 8-(piperidin-4-yl)-2-(2-phenylethyl)-1-methyl-7-benzyl-1,7-dihydro-purin-6-one

(239) 8-(3-amino-hexahydroazepin-1-yl)-2-(2-phenylethyl)-1-methyl-7-benzyl-1,7-dihydro-purin-6-one

(240) 8-(4-amino-hexahydroazepin-1-yl)-2-(2-phenylethyl)-1-methyl-7-benzyl-1,7-dihydro-purin-6-one

(241) 8-(piperazin-1-yl)-2-(2-phenylethyl)-1-methyl-7-benzyl-1,7-dihydro-purin-6-one

(242) 8-(1,4-diazepan-1-yl)-2-(2-phenylethyl)-1-methyl-7-benzyl-1,7-dihydro-purin-6-one

(243) 8-(2-amino-cyclohexylamino)-2-(2-phenylethyl)-1-methyl-7-benzyl-1,7-dihydro-purin-6-one

(244) 8-(3-amino-cyclohexyl)-2-(2-phenylethyl)-1-methyl-7-benzyl-1,7-dihydro-purin-6-one

(245) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(but-2-en-1-yl)-1,7-dihydro-purin-6-one

(246) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(2-methyl-but-2-en-1-yl)-1,7-dihydro-purin-6-one

(247) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(2,3-dimethyl-but-2-en-1-yl)-1,7-dihydro-purin-6-one

(248) 8-(3-amino-piperidin-1-yl)-2-benzylamino-1-methyl-7-(cyclopent-1-en-1-yl-methyl)-1,7-dihydro-purin-6-one

(249) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2-phenyl-2-oxo-ethyl)-1,7-dihydro-purin-6-one

(250) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(2-amino-phenyl)-2-oxo-ethyl]-1,7-dihydro-purin-6-one

(251) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(2-acetylamino-phenyl)-2-oxo-ethyl]-1,7-dihydro-purin-6-one

(252) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(2-methylsulphonylamino-phenyl)-2-oxo-ethyl]-1,7-dihydro-purin-6-one

(253) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(2-hydroxy-phenyl)-2-oxo-ethyl]-1,7-dihydro-purin-6-one

(254) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-[2-(2-methoxy-phenyl)-2-oxo-ethyl]-1,7-dihydro-purin-6-one

(255) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-{2-[2-(ethoxycarbonylmethoxy)-phenyl]-2-oxo-ethyl}-1,7-dihydro-purin-6-one

(256) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-{2-[2-(aminocarbonylmethoxy)-phenyl]-2-oxo-ethyl}-1,7-dihydro-purin-6-one

(257) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-{2-[2-(methylaminocarbonylmethoxy)-phenyl]-2-oxo-ethyl}-1,7-dihydro-purin-6-one

(258) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-{2-[2-(ethylaminocarbonylmethoxy)-phenyl]-2-oxo-ethyl}-1,7-dihydro-purin-6-one

- (259) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-{2-[2-(dimethylaminocarbonyl-methoxy)-phenyl]-2-oxo-ethyl}-1,7-dihydro-purin-6-one
- (260) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(naphth-1-yl-methyl)-1,7-dihydro-purin-6-one
- (261) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(isoquinolin-1-yl-methyl)-1,7-dihydro-purin-6-one
- (262) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(quinazolin-2-yl-methyl)-1,7-dihydro-purin-6-one
- (263) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(quinolin-4-yl-methyl)-1,7-dihydro-purin-6-one
- (264) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(quinazolin-4-yl-methyl)-1,7-dihydro-purin-6-one
- (265) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(2-phenylethyl)-1,7-dihydro-purin-6-one
- (266) 8-[(2-aminoethyl)amino]-7-benzyl-1-methyl-2-benzylamino-1,7-dihydro-purin-6-one
- (267) 8-[N-methyl-N-(2-aminoethyl)-amino]-7-benzyl-1-methyl-2-benzylamino-1,7-dihydro-purin-6-one
- (268) 8-[N-ethyl-N-(2-aminoethyl)-amino]-7-benzyl-1-methyl-2-benzylamino-1,7-dihydro-purin-6-one
- (269) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(2-cyanophenyl)ethyl]-1,7-dihydro-purin-6-one

- (270) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(2-aminocarbonyl-phenyl)ethyl]-1,7-dihydro-purin-6-one
- (271) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(2-dimethylaminocarbonyl-phenyl)ethyl]-1,7-dihydro-purin-6-one
- (272) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(2-methylsulphonyl-phenyl)ethyl]-1,7-dihydro-purin-6-one
- (273) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[2-(2-methylsulphonylamino-phenyl)ethyl]-1,7-dihydro-purin-6-one
- (274) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(2-methylsulphonylamino-ethyl)amino]-1,7-dihydro-purin-6-one
- (275) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(2-phenylsulphonylamino-ethyl)amino]-1,7-dihydro-purin-6-one
- (276) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(naphth-1-yl-methyl)amino]-1,7-dihydro-purin-6-one
- (277) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(quinazolin-2-yl-methyl)amino]-1,7-dihydro-purin-6-one
- (278) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(isoquinolin-1-yl-methyl)-amino]-1,7-dihydro-purin-6-one
- (279) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(4,5-dimethyl-oxazol-2-yl-methyl)-1,7-dihydro-purin-6-one
- (280) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(4,5-dimethyl-thiazol-2-yl-methyl)-1,7-dihydro-purin-6-one

(281) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(4,6-dimethyl-pyrimidin-2-yl-methyl)-1,7-dihydro-purin-6-one

(282) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-(pyrazin-2-yl-methyl)-1,7-dihydro-purin-6-one

(283) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(3-methyl-isoxazol-5-yl-methyl)-amino]-1,7-dihydro-purin-6-one

(284) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(4,5-dimethyl-oxazol-2-yl-methyl)-amino]-1,7-dihydro-purin-6-one

(285) 8-(3-amino-piperidin-1-yl)-7-benzyl-1-methyl-2-[(pyrazin-2-yl-methyl)-amino]-1,7-dihydro-purin-6-one

Example 2

Coated tablets containing 75 mg of active substance

1 tablet core contains:

active substance	75.0 mg
calcium phosphate	93.0 mg
corn starch	35.5 mg
polyvinylpyrrolidone	10.0 mg
hydroxypropylmethylcellulose	15.0 mg
magnesium stearate	<u>1.5 mg</u>
	230.0 mg

Preparation:

The active substance is mixed with calcium phosphate, corn starch, polyvinylpyrrolidone, hydroxypropylmethylcellulose and half the specified amount of magnesium stearate. Blanks 13 mm in diameter are produced in a tablet-making machine and these are then rubbed through a screen with a mesh size of 1.5 mm using a suitable machine and mixed with the rest of the magnesium

stearate. This granulate is compressed in a tablet-making machine to form tablets of the desired shape.

Weight of core: 230 mg

die: 9 mm, convex

The tablet cores thus produced are coated with a film consisting essentially of hydroxypropylmethylcellulose. The finished film-coated tablets are polished with beeswax.

Weight of coated tablet: 245 mg.

Example 3Tablets containing 100 mg of active substance

Composition:

1 tablet contains:

active substance	100.0 mg
lactose	80.0 mg
corn starch	34.0 mg
polyvinylpyrrolidone	4.0 mg
magnesium stearate	<u>2.0 mg</u>
	220.0 mg

Method of Preparation:

The active substance, lactose and starch are mixed together and uniformly moistened with an aqueous solution of the polyvinylpyrrolidone. After the moist composition has been screened (2.0 mm mesh size) and dried in a rack-type drier at 50°C it is screened again (1.5 mm mesh size) and the lubricant is added. The finished mixture is compressed to form tablets.

Weight of tablet: 220 mg

Diameter: 10 mm, biplanar, faceted on both sides and notched on one side.

Example 4Tablets containing 150 mg of active substance

Composition:

1 tablet contains:

active substance	150.0 mg
powdered lactose	89.0 mg
corn starch	40.0 mg
colloidal silica	10.0 mg
polyvinylpyrrolidone	10.0 mg
magnesium stearate	<u>1.0 mg</u>
	300.0 mg

Preparation:

The active substance mixed with lactose, corn starch and silica is moistened with a 20% aqueous polyvinylpyrrolidone solution and passed through a screen with a mesh size of 1.5 mm. The granules, dried at 45°C, are passed through the same screen again and mixed with the specified amount of magnesium stearate. Tablets are pressed from the mixture.

Weight of tablet: 300 mg

die: 10 mm, flat

Example 5Hard gelatine capsules containing 150 mg of active substance

1 capsule contains:

active substance	150.0 mg
corn starch (dried)	approx. 80.0 mg
lactose (powdered)	approx. 87.0 mg
magnesium stearate	3.0 mg
	<hr/>
	approx. 420.0 mg

Preparation:

The active substance is mixed with the excipients, passed through a screen with a mesh size of 0.75 mm and homogeneously mixed using a suitable apparatus. The finished mixture is packed into size 1 hard gelatine capsules.

Capsule filling: approx. 320 mg

Capsule shell: size 1 hard gelatine capsule.

Example 6Suppositories containing 150 mg of active substance

1 suppository contains:

active substance	150.0 mg
polyethyleneglycol 1500	550.0 mg
polyethyleneglycol 6000	460.0 mg
polyoxyethylene sorbitan monostearate	<u>840.0 mg</u>
	2,000.0 mg

Preparation:

After the suppository mass has been melted the active substance is homogeneously distributed therein and the melt is poured into chilled moulds.

Example 7Suspension containing 50 mg of active substance

100 ml of suspension contain:

active substance	1.00 g
carboxymethylcellulose-Na-salt	0.10 g
methyl p-hydroxybenzoate	0.05 g
propyl p-hydroxybenzoate	0.01 g
glucose	10.00 g
glycerol	5.00 g
70% sorbitol solution	20.00 g
flavouring	0.30 g
dist. water	ad 100 ml

Preparation:

The distilled water is heated to 70°C. The methyl and propyl p-hydroxybenzoates together with the glycerol and sodium salt of carboxymethylcellulose are dissolved therein with stirring. The solution is cooled to ambient temperature and the active substance is added and homogeneously dispersed therein with stirring. After the sugar, the sorbitol

solution and the flavouring have been added and dissolved, the suspension is evacuated with stirring to eliminate air.

5 ml of suspension contain 50 mg of active substance.

Example 8

Ampoules containing 10 mg active substance

Composition:

active substance	10.0 mg
0.01 N hydrochloric acid q.s.	
double-distilled water	ad 2.0 ml

Preparation:

The active substance is dissolved in the necessary amount of 0.01 N HCl, made isotonic with common salt, filtered sterile and transferred into 2 ml ampoules.

Example 9

Ampoules containing 50 mg of active substance

Composition:

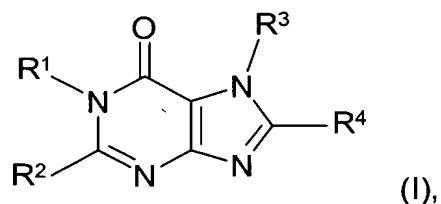
active substance	50.0 mg
0.01 N hydrochloric acid q.s.	
double-distilled water	ad 10.0 ml

Preparation:

The active substance is dissolved in the necessary amount of 0.01 N HCl, made isotonic with common salt, filtered sterile and transferred into 10 ml ampoules.

Abstract

The invention relates to new purine derivatives of general formula



wherein R^1 to R^4 are defined as in the claims, the tautomers, the stereoisomers, the mixtures, the prodrugs thereof and the salts thereof, particularly the physiologically acceptable salts thereof with inorganic or organic acids or bases which have valuable pharmacological properties, particularly an inhibiting effect on the activity of the enzyme dipeptidyl-peptidase-IV (DPP-IV), the preparation thereof, the use thereof for the prevention or treatment of diseases or conditions associated with an increased DPP-IV activity or capable of being prevented or alleviated by reducing the DPP-IV activity, particularly type I or type II diabetes mellitus, the pharmaceutical compositions containing a compound of general formula (I) or a physiologically acceptable salt thereof as well as processes for the preparation thereof.